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(54) **PAIN ASSESSMENT**

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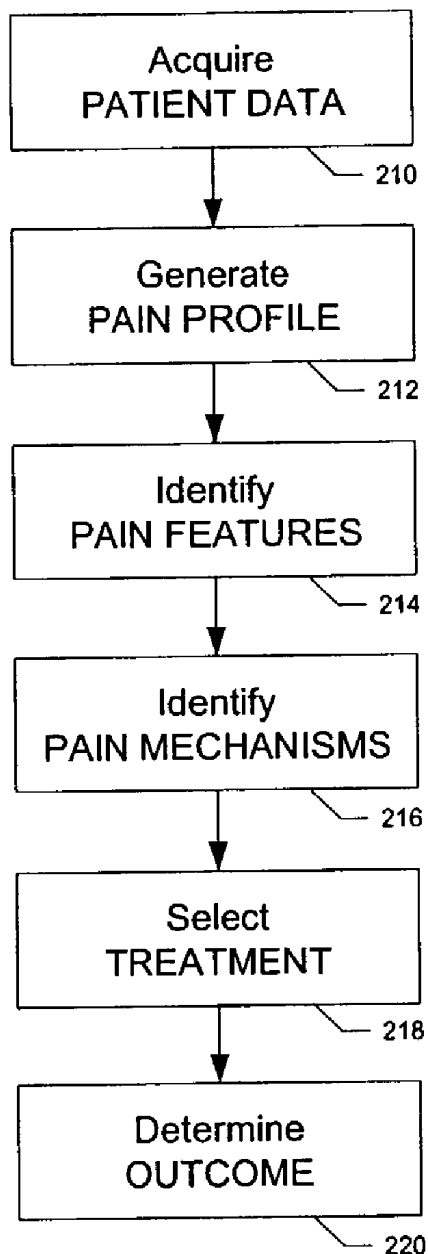
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(57) **ABSTRACT**

Methods and questionnaires are provided for assessing pain.

**START**



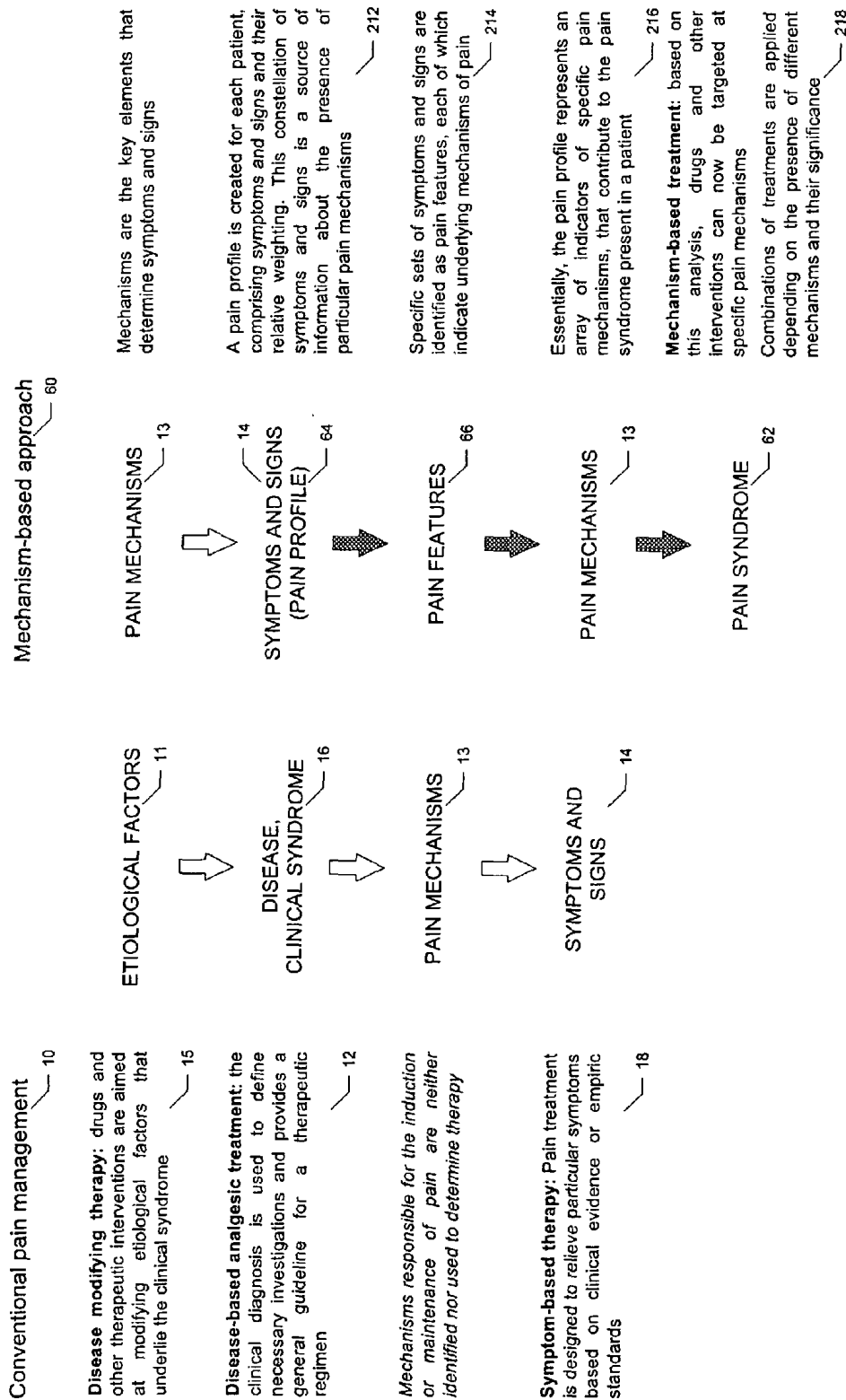


Figure 1

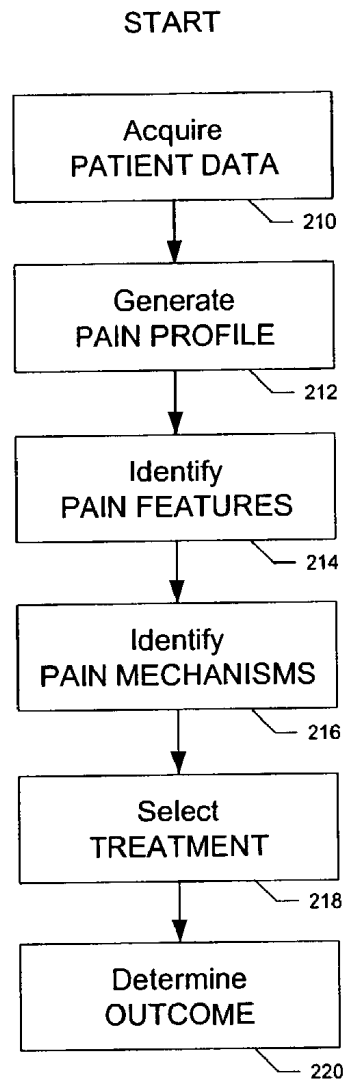


Figure 1A

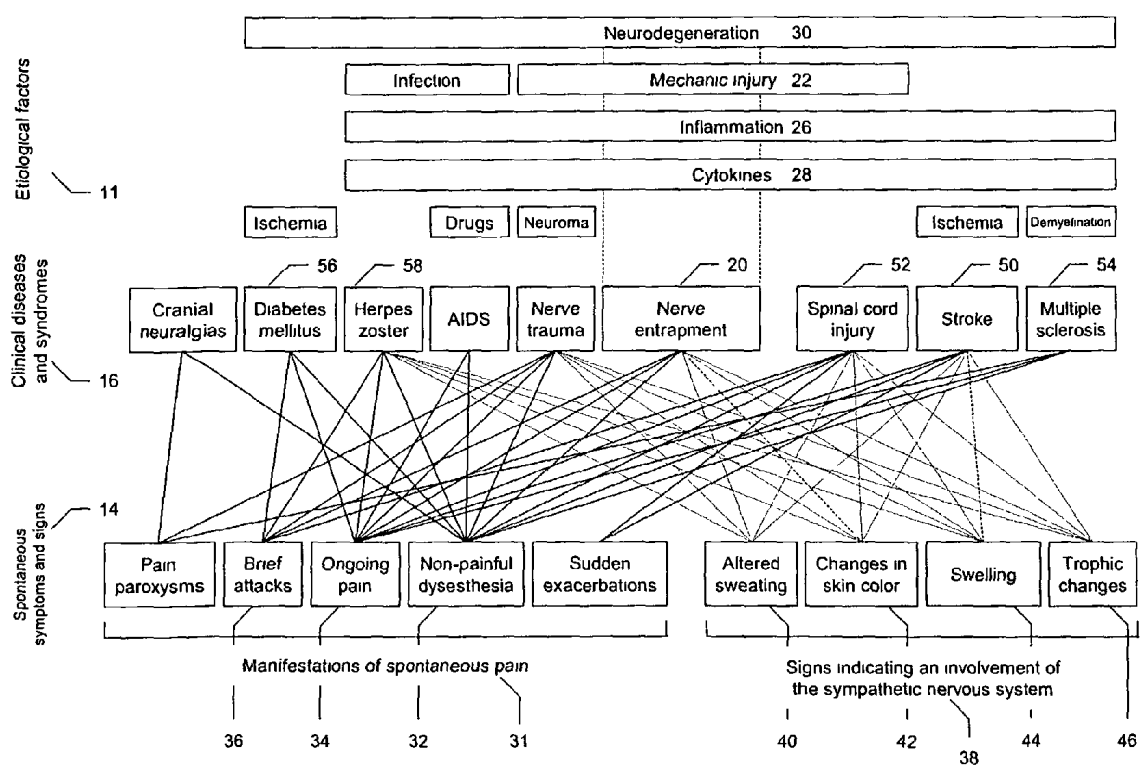


Figure 2

## Mechanisms underlying symptoms and signs in conditions of neuropathic pain

| Spontaneous sensory disorders                                                                                    |                                                                                   | Mechanisms                                                                                                                                                                                                                               |                                                                                                                      | Stimulus-evoked sensory disorders |                                                                                              |
|------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------|-----------------------------------|----------------------------------------------------------------------------------------------|
| A $\beta$ -nerve fibers                                                                                          | A $\delta$ or C-nerve fibers                                                      | Peripheral nervous system (Sensory system)                                                                                                                                                                                               | A $\beta$ -nerve fibers                                                                                              | A $\delta$ or C-nerve fibers      |                                                                                              |
| Decrease or complete loss of the sensation of touch (static or dynamic), position and movement, and vibration    | Decrease or complete loss of the sensation of cold, warm (hot), and pinprick pain | Degeneration of the peripheral sensory nerve fiber terminal, the peripheral or the central axon branch, the dorsal root ganglion neurons, the central sensory nerve fiber terminal in the dorsal horn, or ascending dorsal column fibers | N/A                                                                                                                  | N/A                               |                                                                                              |
|                                                                                                                  | Painful (burning) sensation                                                       | Reduced threshold of activation in nociceptors of the peripheral terminal                                                                                                                                                                |                                                                                                                      |                                   | Pinprick hyperalgesia in A $\delta$ fibers, heat hyperalgesia and cold allodynia in C-fibers |
| Paresthesia, dysesthesia                                                                                         | Painful (burning) sensation                                                       | Increased excitability along the axon, in the dorsal root ganglion neuron or the dorsal root                                                                                                                                             |                                                                                                                      |                                   | Pain caused by stretching (straight-leg-raising sign)                                        |
|                                                                                                                  |                                                                                   | Phenotypic changes with the increased or novel expression of receptors and ion channels along the axon, or in the dorsal root ganglion neuron                                                                                            | Paresthesia, dysesthesia on direct pressure (Tinel sign)                                                             |                                   | Painful neuroma sign                                                                         |
| Paresthesia, dysesthesia                                                                                         | Painful (burning) sensation                                                       | Ectopic activity along the axon, in the dorsal root ganglion neuron or the dorsal root                                                                                                                                                   |                                                                                                                      |                                   |                                                                                              |
|                                                                                                                  |                                                                                   | Central nervous system (Processing the sensory input)                                                                                                                                                                                    |                                                                                                                      |                                   |                                                                                              |
|                                                                                                                  |                                                                                   | Abnormal sensory excitability due to an increased synaptic release of neurotransmitter or enhanced post-synaptic sensitivity                                                                                                             | Allodynia on touch (static or dynamic), pressure, movement                                                           |                                   | Hyperalgesia on stimulation with pinprick, heat, or cold                                     |
| Diffuse pain (accompanied by mood changes, sleep disturbance)                                                    |                                                                                   | Widespread phenotypic changes of neurons in the central nervous system, e.g. induced by expression of COX-2 during inflammation                                                                                                          |                                                                                                                      |                                   | Diffuse hyperalgesia                                                                         |
|                                                                                                                  |                                                                                   | Structural reorganization of synapses                                                                                                                                                                                                    | Allodynia on touch (static or dynamic) (especially within an area of reduced or lost pinprick and thermal sensation) |                                   |                                                                                              |
| Paresthesia, dysesthesia and pain                                                                                |                                                                                   | Loss of inhibition                                                                                                                                                                                                                       | Allodynia on touch, pressure, movement, cold                                                                         |                                   | Hyperalgesia on stimulation with pinprick, heat, or cold                                     |
| Complex sensation in a distribution exceeding the boundaries of dorsal root or peripheral nerve-innervated areas |                                                                                   | Spontaneous activity in neurons of the dorsal horn or higher centers, due to increased excitability or decreased inhibition, or both                                                                                                     |                                                                                                                      |                                   |                                                                                              |
|                                                                                                                  |                                                                                   | Sympathetic nervous system (Central and peripheral)                                                                                                                                                                                      |                                                                                                                      |                                   |                                                                                              |
| Altered sweating, changes in skin blood flow, edema                                                              |                                                                                   | Altered level of activity                                                                                                                                                                                                                |                                                                                                                      |                                   |                                                                                              |
| Paresthesia, dysesthesia or painful burning sensation                                                            |                                                                                   | Sympathetic-sensory coupling                                                                                                                                                                                                             |                                                                                                                      |                                   |                                                                                              |

Figure 3

Two examples of the relationship between pain features and composite symptoms and signs

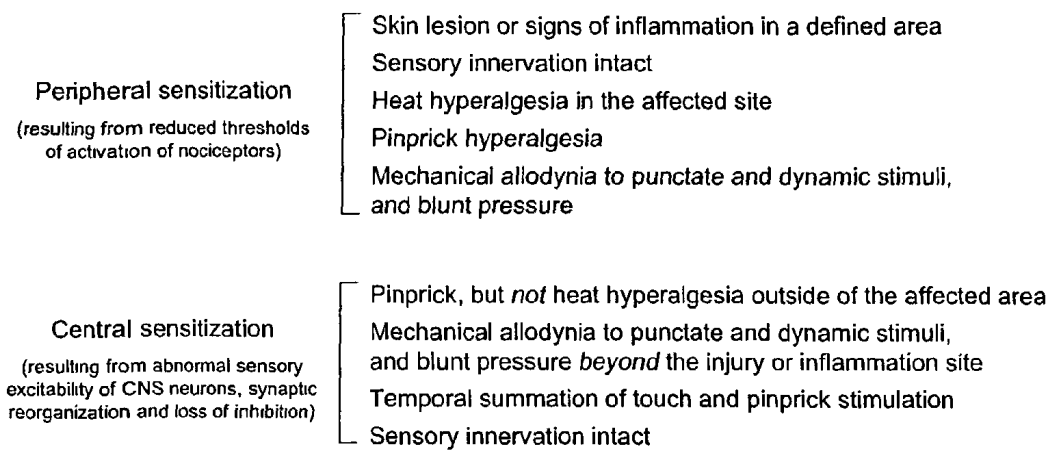


Figure 4

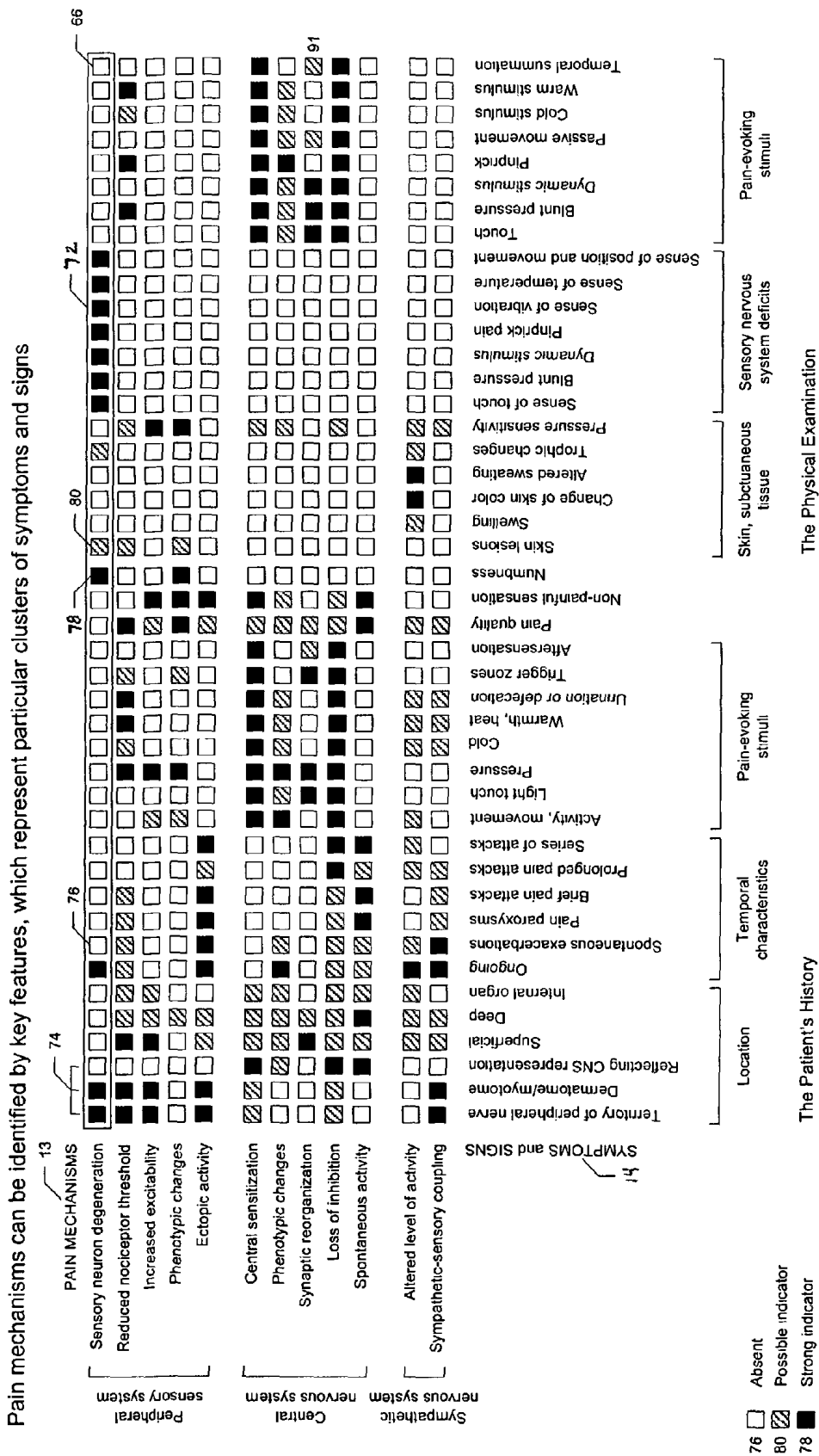


Figure 5

**Patient**  
*Clinical diagnosis: Postherpetic neuralgia*

| The Patient's History |                  | Pain 1          |  | Pain 2 |  |
|-----------------------|------------------|-----------------|--|--------|--|
| Location              | Trunk (T5, left) | Same            |  |        |  |
| Superficial/deep      | Deep             | Superficial     |  |        |  |
| Ongoing/intermittent  | Ongoing          | Evoked by touch |  |        |  |
| Spontaneous/evoked    | Spontaneous      | NRS 6           |  |        |  |
| Intensity             | NRS 5            | Aching          |  |        |  |
| Quality               | Severe           | Sharp           |  |        |  |
| Numeness              |                  |                 |  |        |  |

| The Physical Examination    |                     |
|-----------------------------|---------------------|
| Skin lesions                | Scars               |
| Sensitivity of deep tissues | Testing evokes pain |
| Sense of touch              | Mild decrease       |
| Pinprick intensity          | Severe decrease     |
| Sense of vibration          | Mild decrease       |
| Sense of temperature        | Severe decrease     |
| Intensity of evoked pain    |                     |
| Pain evoking stimuli        | Mechanical          |

**The Patient's History**

Location: ☐ Superficial ☐ Deep ☐ Internal organ ☐ Ongoing ☐ Pain paroxysms ☐ Brief pain attacks ☐ Prolonged pain attacks ☐ Series of attacks ☐ Activity, movement ☐ Light touch ☐ Pressure ☐ Cold ☐ Warmth, heat ☐ Urination or defecation ☐ Trigger zones ☐ Allodynia ☐ Pain quality ☐ Non-painful sensation ☐ Numbness ☐ Skin lesions ☐ Swelling ☐ Change of skin color ☐ Altered sweating ☐ Trophic changes ☐ Pressure sensitivity ☐ Sense of touch ☐ Blunt pressure ☐ Dynamic stimulus ☐ Pinprick pain ☐ Sense of vibration ☐ Sense of temperature ☐ Sense of position and movement ☐ Touch ☐ Blunt pressure ☐ Dynamic stimulus ☐ Pinprick ☐ Passive movement ☐ Cold stimulus ☐ Warm stimulus ☐ Temporal summation

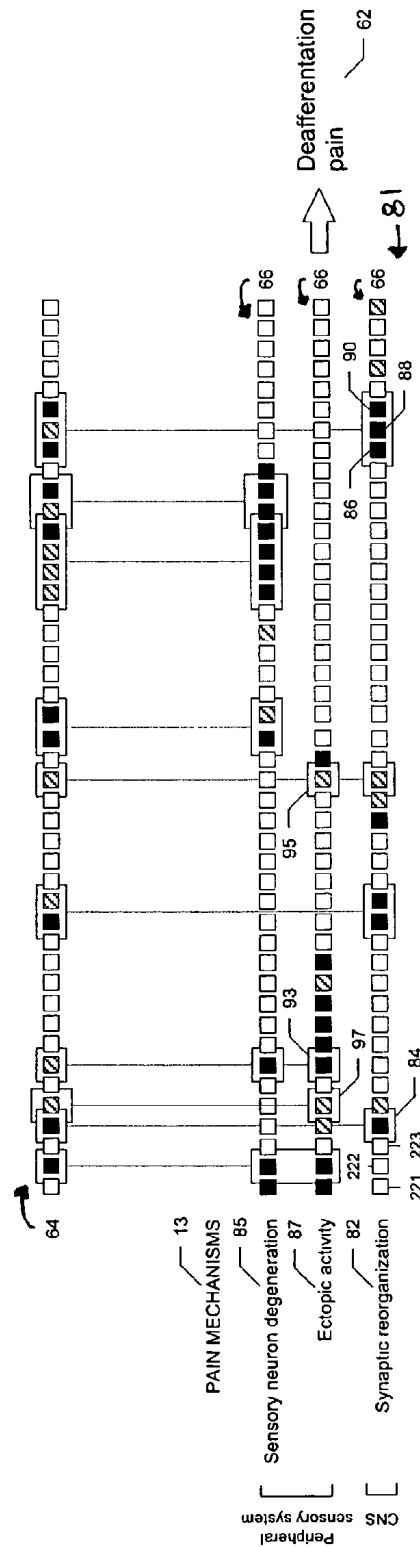
**The Physical Examination**

Skin, subcutaneous tissue: ☐ Sensory system deficits ☐ Pain-evoking stimuli

**Legend:** ☐ Absent ☐ Low rating ☐ High rating

**STEP 1** Symptoms and signs are investigated using a standardized questionnaire

**STEP 2** Symptoms and signs create an individual patient's pain profile



**STEP 3** Particular pain features allow to identify operating pain mechanisms. Their combination provides a mechanism-based diagnosis of a pain syndrome

**STEP 4**

Figure 6

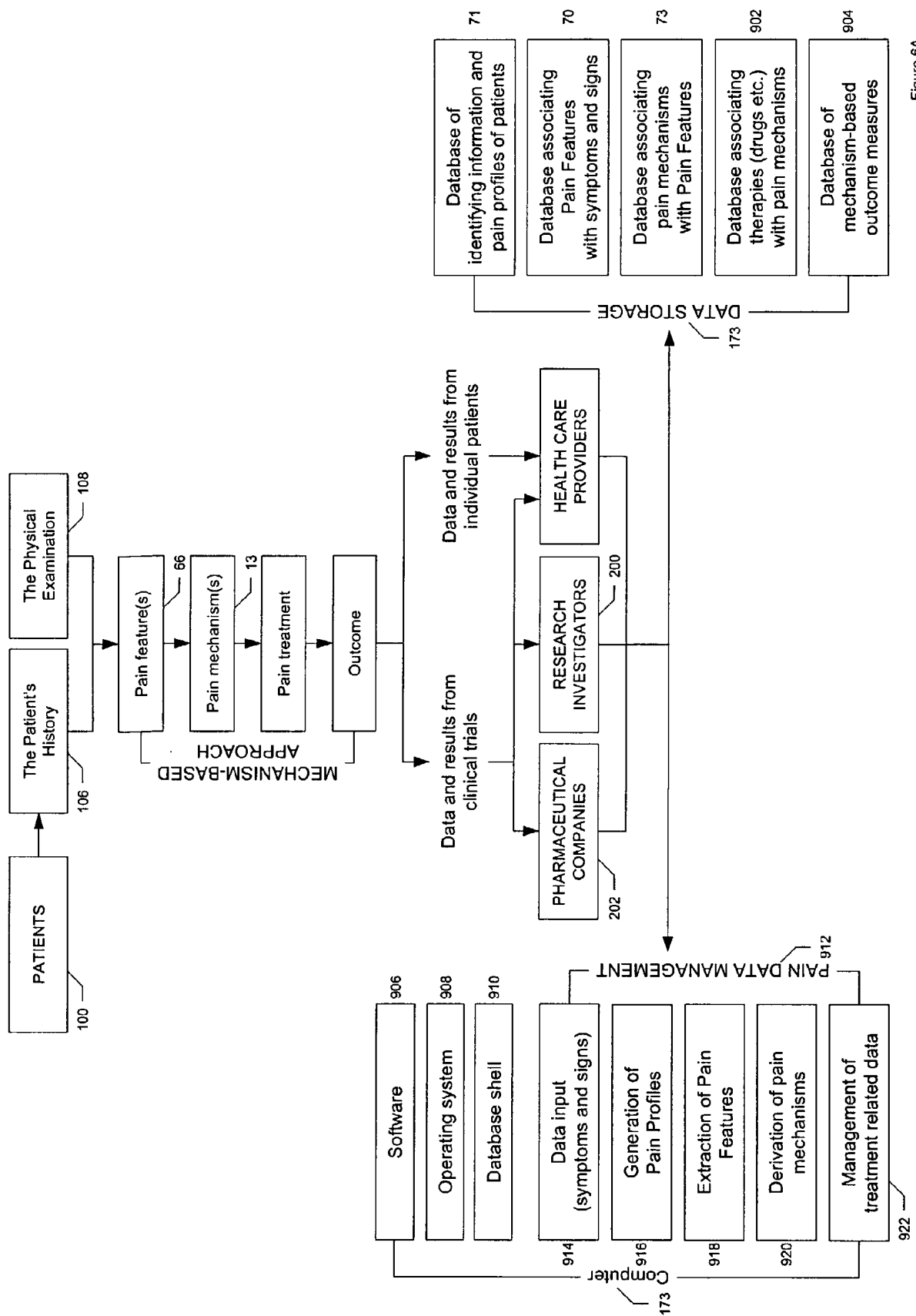


Figure 6A

## Mechanisms-based treatment options for neuropathic pain

180

| Mechanism                                            | Target                                                                                                                                              | Drug treatment                                                                                                                                                                                                                                                                                                                                                                                   |
|------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Peripheral nervous system</b><br>(Sensory system) |                                                                                                                                                     |                                                                                                                                                                                                                                                                                                                                                                                                  |
| Degeneration of the peripheral sensory neuron        | Neurotrophin-receptors<br>Caspases                                                                                                                  | Neurotrophins<br>Inhibitors of caspase activation (Hsp27)<br>Caspase inhibitors (zVAD.fmk, DEVD.fmk)                                                                                                                                                                                                                                                                                             |
| Reduced threshold of activation of nociceptors       | VR1<br>Bradykinin<br>Serotonin<br>NGF                                                                                                               | Capsaicin<br>VR1 antagonists<br>Bradykinin antagonists<br>Serotonin antagonists<br>NGF antagonists                                                                                                                                                                                                                                                                                               |
| Peripheral sensory neuron hyperexcitability          | TTXr-VGSC<br>Potassium channels                                                                                                                     | Selective blockers of TTXr-VGSC<br>Potassium channel activators                                                                                                                                                                                                                                                                                                                                  |
| Phenotypic changes                                   | Substance P                                                                                                                                         | Substance P antagonists                                                                                                                                                                                                                                                                                                                                                                          |
| Ectopic activity                                     | TTXs-VGSC<br>TTXr-VGSC<br>Potassium channels                                                                                                        | Sodium channel blockers (local anesthetics, carbamazepine, lamotrigine, mexiletine)<br>Selective blockers of TTXr-VGSC<br>Potassium channel activators                                                                                                                                                                                                                                           |
| <b>Central nervous system</b>                        |                                                                                                                                                     |                                                                                                                                                                                                                                                                                                                                                                                                  |
| Abnormal sensory excitability                        | NMDA-R<br><br>Kainate receptor<br>mGlu-R<br>NK1<br>nNOS<br>PKC $\gamma$<br>MAPK/ERK<br>N-type Ca <sup>2+</sup> channels<br>VGCC, $\alpha$ 2-subunit | NMDA-R antagonists (ketamine, dextromethorphan, amantadine)<br>Glycine site antagonists<br>NR2A subunit antagonists<br>Kainate receptor antagonists<br>mGlu-R antagonists<br>NK1 antagonists<br>nNOS inhibitors<br>PKC $\gamma$ inhibitors<br>MAPK/ERK inhibitors<br>$\omega$ -Conotoxin<br>Gabapentin                                                                                           |
| Phenotypic changes                                   | COX-2                                                                                                                                               | COX-2 inhibitors                                                                                                                                                                                                                                                                                                                                                                                 |
| Synaptic reorganization                              | GFR $\alpha$ -1(2)/RET                                                                                                                              | GDNF                                                                                                                                                                                                                                                                                                                                                                                             |
| Loss of inhibition                                   | MOR<br>CB-2<br>GABA<br>$\alpha$ 2-Adenoreceptor<br><br>Adenosine receptor<br>P2X <sub>3</sub><br>CCK<br>nAch-R                                      | $\mu$ -Opioid agonists (morphine, oxycodone)<br>CB-2 agonists (cannabinoid)<br>GABA <sub>A</sub> , GABA <sub>B</sub> agonists (baclophen)<br>$\alpha$ 2-Adenoreceptor agonists (clonidine)<br>Tricyclic antidepressants (Amitriptyline, Imipramine, Clomipramine, desipramine, maprotiline)<br>Adenosine receptor agonists<br>P2X <sub>3</sub> antagonists<br>CCK antagonists<br>nAch-R agonists |
| Spontaneous activity                                 | N-type Ca <sup>2+</sup> channels<br>VGCC, $\alpha$ 2-subunit                                                                                        | $\omega$ -Conotoxin<br>Gabapentin                                                                                                                                                                                                                                                                                                                                                                |
| <b>Sympathetic nervous system</b>                    |                                                                                                                                                     |                                                                                                                                                                                                                                                                                                                                                                                                  |
| Altered level of activity                            | $\alpha$ 2-Adenoreceptor<br>VGSC                                                                                                                    | $\alpha$ 2-Adenoreceptor agonists (clonidine)<br>Sympathetic blocks using local anesthetics                                                                                                                                                                                                                                                                                                      |
| Sympathetic-sensory coupling                         | $\alpha$ 1-Adenoreceptor<br>NGF<br>VGSC                                                                                                             | $\alpha$ 1-Adenoreceptor agonists (phenolamine, guanethidine)<br>NGF antagonists<br>Sympathetic blocks using local anesthetics                                                                                                                                                                                                                                                                   |

Figure 7

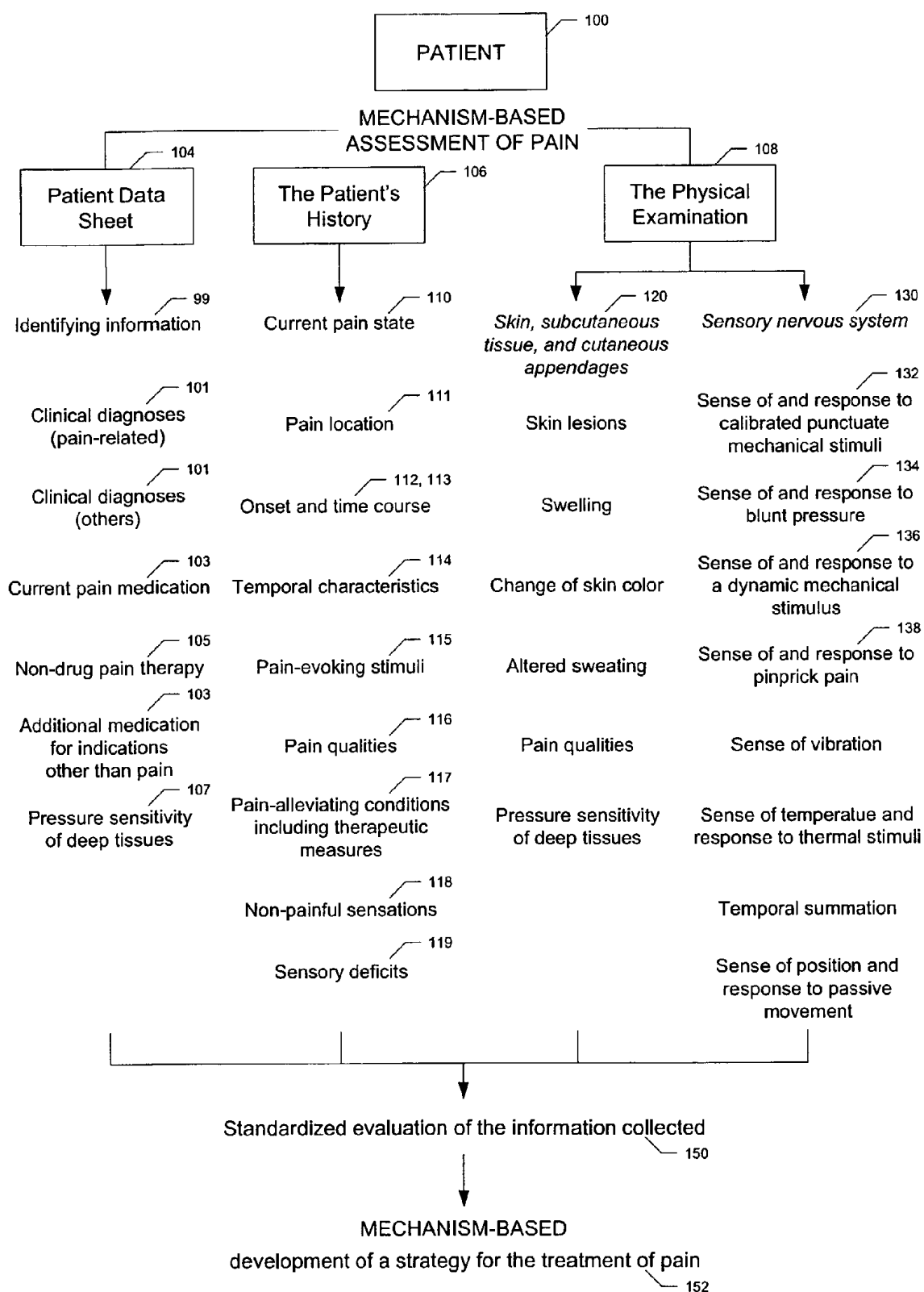


Figure 8

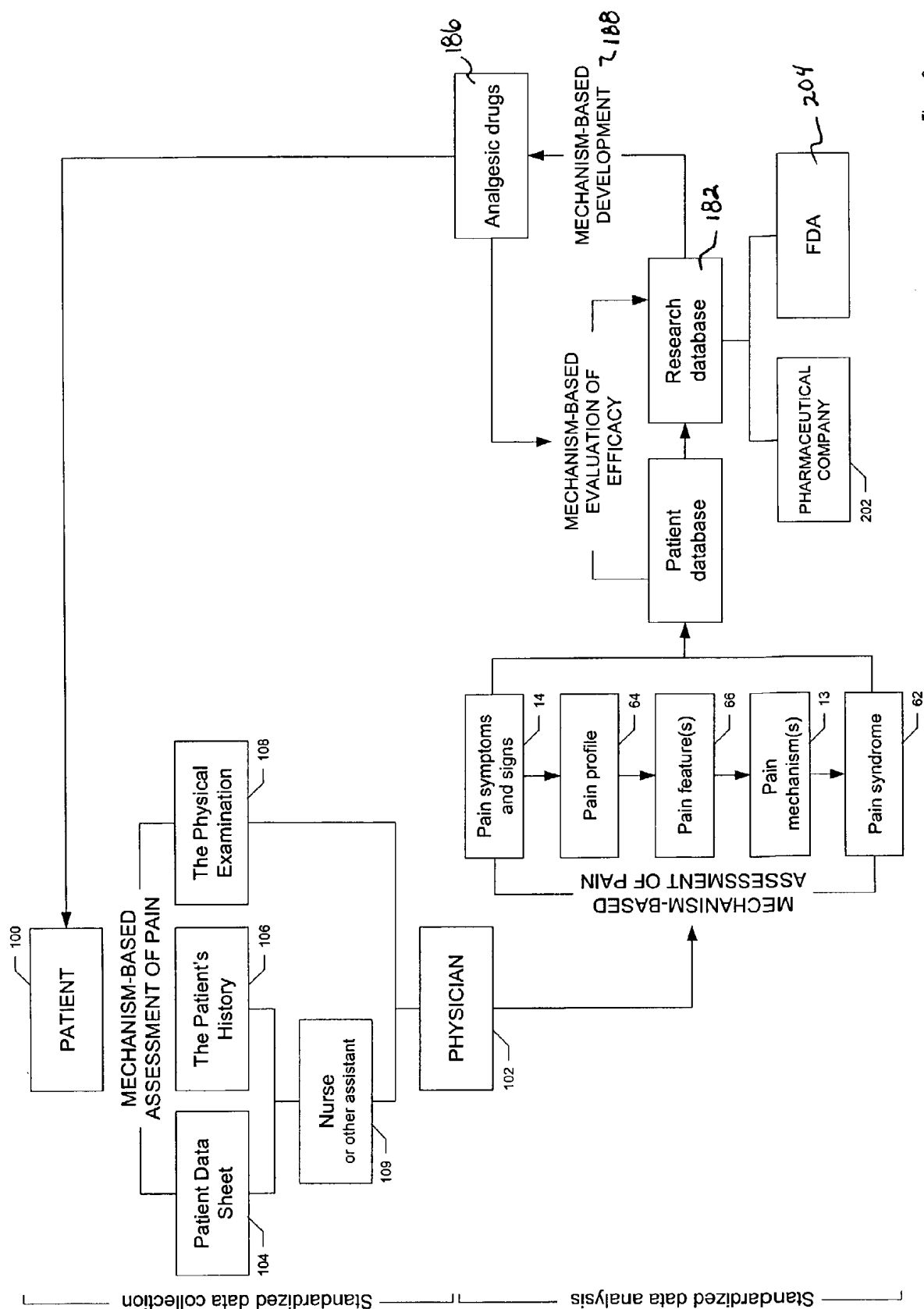


Figure 9

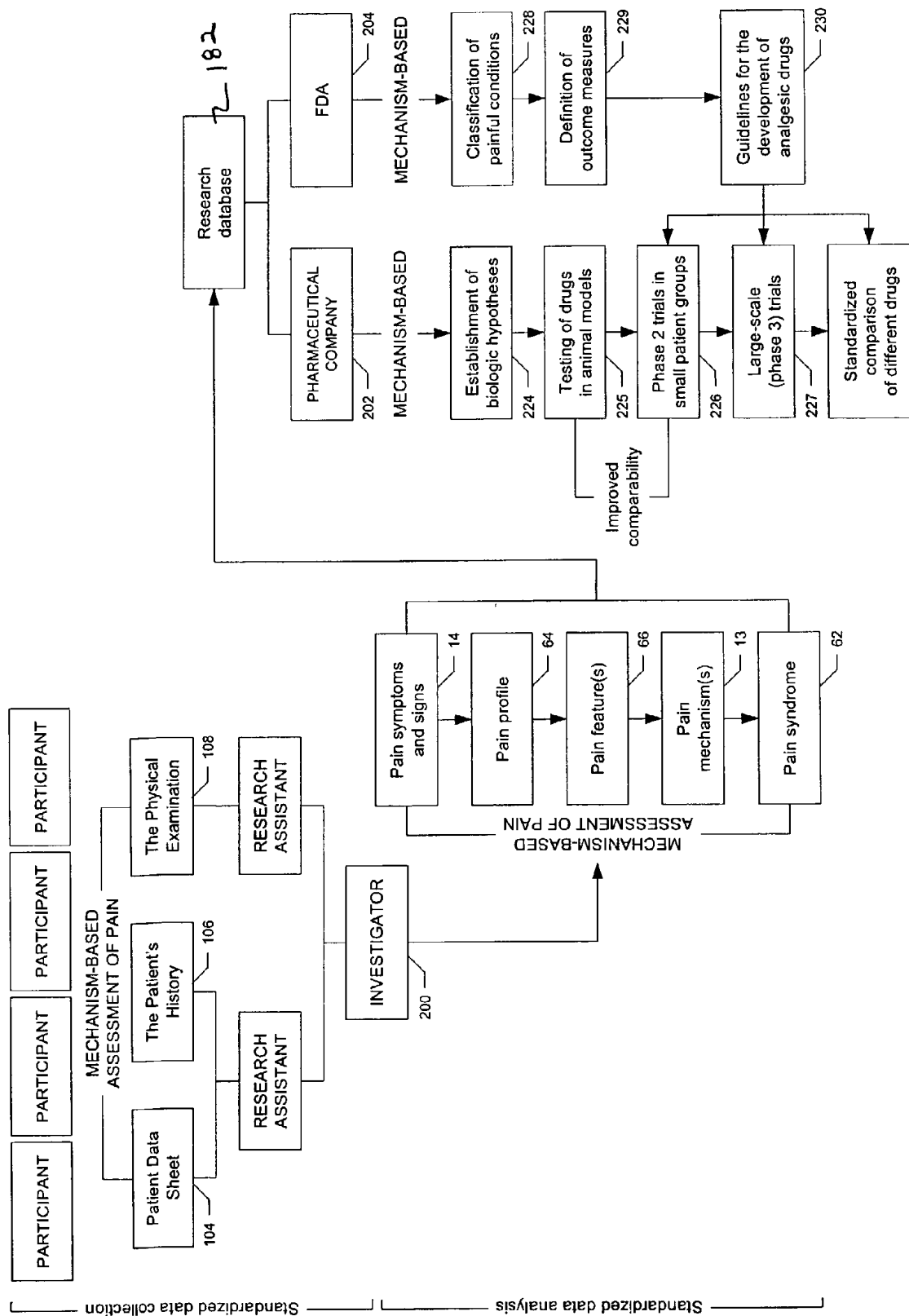


Figure 10

## ***Clinical Assessment of Pain Syndromes***

### ***The Patient's History***

Date ..... / ..... / .....  
(MM/DD/YY)

Patient's First Name .....  
Last Name .....  
Hospital/Patient Number .....  
Date of Birth ..... / ..... / .....  
☐ Female      ☐ Male

Investigator's Name .....

*Ask the patient the following questions and enter the responses*

**Current Pain State**

**Section H1**

H1-01 **Are you in pain right now?**

- ☐ No  
☐ Yes

Describe the pain location briefly, e.g. "right hand" .....

Rate the **intensity of the current pain** by checking one of the four descriptors

- ☐ none      ☐ mild      ☐ moderate      ☐ severe

**and**

on a numerical scale from 0 to 10  where 0 represents no pain and 10 the maximum possible pain

**Pain Location**

**Section H2**

H2-01 **Where is the pain located that you consider to be usually the most disturbing or distressing?**

Describe the location briefly .....

**and**

mark the whole affected area on the drawing on the following page. Assign number **1** for this pain in the drawing

If there is a clear spot or a defined area where this pain is most pronounced or extreme, label it with a cross, "x"

H2-02 **Is there another (or more than one other) pain that differs either in the type of sensation, the location or the frequency from Pain No. 1?**

- ☐ No  
☐ Yes

Describe the location of this second (or additional) pain(s) briefly .....

**and**

mark it (or them) on the drawing on the following page. Assign number **2** for the second pain in the drawing. If there are more than two different types of pain, use successive numbers

If there is a clear spot or a defined area where this second or any additional pain is most pronounced or extreme, label it with a cross, "x"

*In the following, answers need to be assigned to Pain No. 1 and/or Pain No. 2. If there are more than two types of pain, use a separate sheet and answer the questions correspondingly*

H2-03 **Is your pain superficial, deep, or felt as if in an internal organ?** —————→ Pain No. 1 Pain No. 2

- Superficial (on or in the skin) ☐ ☐
  - Deep (under the skin, in muscles, bone, joints etc.) ☐ ☐
  - Felt in or as if in an internal organ of the body (as the heart, liver, or intestine) ☐ ☐
- Specify .....

**Onset and Time Course**

**Section H3**

H3-01 **When did your pain start?** —————→ Pain No. 1 Pain No. 2

- < 3 Months ago ☐ ☐
- < 6 Months ago ☐ ☐
- < 1 Year ago ☐ ☐
- > 1 Year ago ☐ ☐

Specify .....

Figure 11B

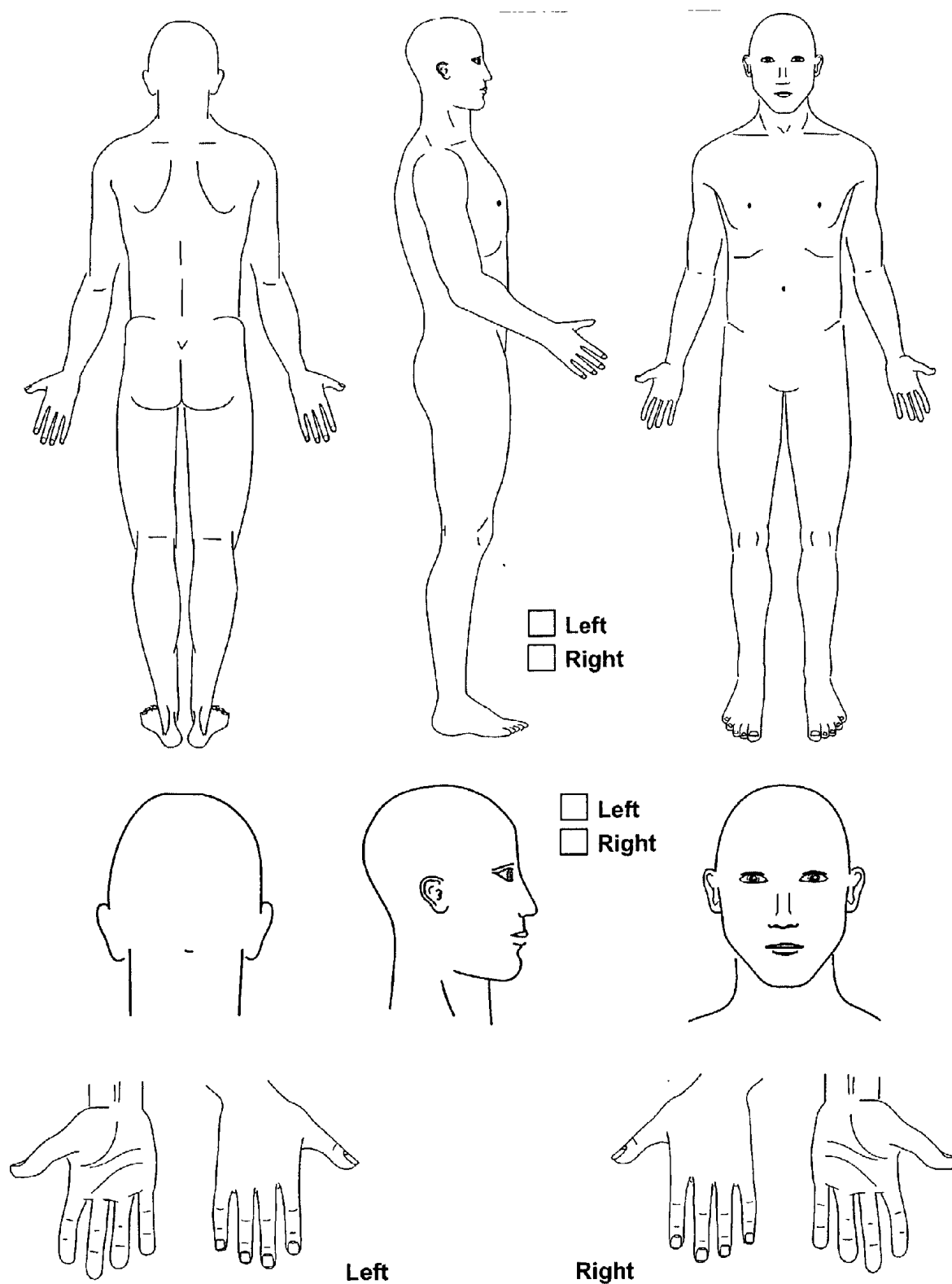


Figure 11C

H3-02 Was there a disease, an injury or any other event that seemed to cause your pain?

☐ No

☐ Yes

- Disease

Specify

- Injury

Specify the type and the site of the injury

- Others

Specify

Pain No. 1

Pain No. 2

☐

☐

☐

☐

☐

☐

☐

☐

H3-03 Has your pain changed over time?

☐ No

☐ Yes

- Pain has decreased in intensity

- Pain has increased in intensity

- Pain has changed its character

Specify the change

Pain No. 1

Pain No. 2

☐

☐

☐

☐

☐

☐

☐

☐

#### Temporal Characteristics

#### Section H4

H4-01 Is your pain always there?

☐ No

If "No" is answered, proceed to item H4-04

☐ Yes

Rate the average intensity of this ongoing pain by checking one of the following four descriptors

☐ none

☐ mild

☐ moderate

☐ severe

and

on a numerical scale from 0 to 10 ☐ where 0 represents no pain and 10 the maximum possible pain

Pain No. 1

Pain No. 2

☐

☐

H4-02 Does your pain ever suddenly, spontaneously worsen?

This question targets only **spontaneous** increases in pain severity. Pain evoked by specific events including external stimulation is investigated in the following section

☐ No

If "No" is answered, proceed to item H5-01

☐ Yes, sudden increases in pain intensity occur spontaneously

Rate the intensity of this suddenly increased pain by checking one of the following four descriptors

☐ none

☐ mild

☐ moderate

☐ severe

and

on a numerical scale from 0 to 10 ☐

Pain No. 1

Pain No. 2

☐

☐

H4-03 How often do you suffer from such increases in pain intensity?

☐ Times on average per ☐ day ☐ week ☐ month

Continue the interview now with the next set of questions. Proceed to item H5-01

Pain No. 1

Pain No. 2

☐

☐

Figure 11D

H4-04 If your pain is not always present, how frequently do you experience spontaneous pain attacks, that do not result from a specific event? → Pain No. 1 Pain No. 2  
☐ Times on average per ☐ day ☐ week ☐ month ☐ ☐

H4-05 How long does a single pain attack usually last? → Pain No. 1 Pain No. 2  
☐ < 1 Minute ☐ ☐  
 or  
☐ Minutes ☐ Hours ☐ ☐

Rate the average intensity of spontaneous pain attacks by checking one of the following four descriptors  
☐ none ☐ mild ☐ moderate ☐ severe

and

on a numerical scale from 0 to 10 ☐ where 0 represents no pain and 10 the maximum possible pain

H4-06 Within which time do pain attacks reach their maximal intensity? → Pain No. 1 Pain No. 2  
☐ < 1 Minute ☐ ☐  
 or  
☐ Minutes ☐ Hours ☐ ☐

H4-07 Are there series of attacks?  
☐ No  
☐ Yes → Pain No. 1 Pain No. 2  
 Specify the average duration of such a series:  
☐ Minutes ☐ Hours ☐ ☐

#### Evoked Pain

#### Section H5

H5-01 Do you suffer from pain that results from a specific event, e.g. a movement, a touch, exposure to cold or warmth?

☐ No

If "No" is answered, proceed to item H6-01

☐ Yes →

- Pain is evoked **only** during or immediately after the stimulus and, once the stimulus ends, tends to wear off or fade away
- Evoked pain clearly outlasts the stimulus and remains at an intense level or increases

Pain No. 1 Pain No. 2  
☐ ☐  
☐ ☐  
☐ ☐

H5-02 Are there defined spots where touch, pressure or other stimuli consistently cause a pain attack?

☐ No

☐ Yes →

Describe the location briefly .....

and

mark the spots in the drawing on page 4. Label them with a dot, "●"

Pain No. 1 Pain No. 2  
☐ ☐

H5-03 What seems to produce your pain most often or most intensely? → Pain No. 1 Pain No. 2  
 Specify ..... ☐ ☐

Figure 11E

| Is your pain in particular caused by ... |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   | Pain No. 1               | Pain No. 2               |
|------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------|--------------------------|
| H5-04                                    | <p>• ... <b>activity</b>, e.g. when you are moving an arm or leg, bending or straightening your back, when you are walking, coughing, or chewing?</p> <p>Specify .....</p> <p>Rate the intensity of the <b>activity-evoked pain</b> by checking one of the four descriptors</p> <p><input type="checkbox"/> none      <input type="checkbox"/> mild      <input type="checkbox"/> moderate      <input type="checkbox"/> severe</p> <p><b>and</b></p> <p>on a numerical scale from 0 to 10 <input type="checkbox"/> where 0 represents no pain and 10 the maximum possible pain</p>                                               | <input type="checkbox"/> | <input type="checkbox"/> |
| H5-05                                    | <p>• <b>Is your pain caused by light touch</b> like from a shirt or a bed sheet?</p> <p>Rate the intensity of the <b>touch-evoked pain</b> by checking one of the four descriptors</p> <p><input type="checkbox"/> none      <input type="checkbox"/> mild      <input type="checkbox"/> moderate      <input type="checkbox"/> severe</p> <p><b>and</b></p> <p>on a numerical scale from 0 to 10 <input type="checkbox"/></p> <p>Is this <b>pain</b> steadily there or does it get worse when the touch stimulus is moving?</p> <p><input type="checkbox"/> Steadily there</p> <p><input type="checkbox"/> Worse when moving</p> | <input type="checkbox"/> | <input type="checkbox"/> |
| H5-06                                    | <p>• <b>Is your pain caused by pressure</b> like from a belt, a bra or shoes, or a firm grip?</p> <p>Specify .....</p> <p>Rate the intensity of the <b>pressure-evoked pain</b> by checking one of the four descriptors</p> <p><input type="checkbox"/> none      <input type="checkbox"/> mild      <input type="checkbox"/> moderate      <input type="checkbox"/> severe</p> <p><b>and</b></p> <p>on a numerical scale from 0 to 10 <input type="checkbox"/></p>                                                                                                                                                               | <input type="checkbox"/> | <input type="checkbox"/> |
| H5-07                                    | <p>• <b>Is your pain caused by cold</b>, e.g. cold air on your skin?</p> <p>Specify pain-evoking cold stimuli .....</p> <p>Rate the intensity of the <b>cold-evoked pain</b> by checking one of the four descriptors</p> <p><input type="checkbox"/> none      <input type="checkbox"/> mild      <input type="checkbox"/> moderate      <input type="checkbox"/> severe</p> <p><b>and</b></p> <p>on a numerical scale from 0 to 10 <input type="checkbox"/></p>                                                                                                                                                                  | <input type="checkbox"/> | <input type="checkbox"/> |
| H5-08                                    | <p>• <b>Is your pain caused by warm</b>, e.g. a warm bath?</p> <p>Specify pain-evoking warm stimuli .....</p> <p>Rate the intensity of the <b>warm-evoked pain</b> by checking one of the four descriptors</p> <p><input type="checkbox"/> none      <input type="checkbox"/> mild      <input type="checkbox"/> moderate      <input type="checkbox"/> severe</p> <p><b>and</b></p> <p>on a numerical scale from 0 to 10 <input type="checkbox"/></p>                                                                                                                                                                            | <input type="checkbox"/> | <input type="checkbox"/> |
| H5-09                                    | <p>• <b>Is your pain caused by urination or defecation?</b></p> <p>Specify .....</p>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              | <input type="checkbox"/> | <input type="checkbox"/> |

Item H5-09 continued on the following page

Figure 11F

Rate the intensity of the **pain evoked by urination or defecation** by checking one of the four descriptors

☐ none ☐ mild ☐ moderate ☐ severe

and

on a numerical scale from 0 to 10

### Pain Qualities

### Section H6

H6-01 How would you describe your pain? →

Pain No. 1 Pain No. 2

Specify

☐ ☐

Check whether the description given by the patient matches one or more of the following categories. The categories may be proposed if the patient cannot describe his pain(s) articulately

- Throbbing or pounding or pulsating
- Shooting or radiating
- Cramping or gripping or squeezing
- Stabbing or sharp
- Aching or dull
- Pricking or painful pins and needles
- Burning or hot
- Others

☐ ☐  
☐ ☐  
☐ ☐  
☐ ☐  
☐ ☐  
☐ ☐  
☐ ☐  
☐ ☐

Specify

### Effect of Drugs and Other Pain Alleviating Conditions

### Section H7

H7-01 Can you name drugs or any other treatment(s) that have relieved your pain?

☐ No

Pain No. 1 Pain No. 2

☐ Yes →

☐ ☐

☐ Drugs

☐ ☐

Specify these drugs and if you remember, the maximal dosage taken

(1)

☐ ☐

(2)

☐ ☐

(3)

☐ ☐

(4)

☐ ☐

(5)

☐ ☐

☐ Nerve blocks, including epidural injections and sympathetic blocks

☐ ☐

Specify

☐ Physical therapy

☐ ☐

Specify

☐ Electrical stimulation of nerves like TENS, or spinal cord stimulation (SCS)

☐ ☐

Specify

☐ Acupuncture

☐ ☐

Specify

☐ Others

☐ ☐

Specify

H7-02 Can you also name drugs or any other treatment(s) that did not relieve your pain?

☐ No

☐ Yes

☐ Drugs

Specify these drugs and if you remember, the maximal dosage taken

(1) .....

(2) .....

(3) .....

(4) .....

(5) .....

☐ Nerve blocks, including epidural injections and sympathetic blocks

Specify .....

☐ Physical therapy

Specify .....

☐ Electrical stimulation of nerves like TENS, or spinal cord stimulation (SCS)

Specify .....

☐ Acupuncture

Specify .....

☐ Others

Specify .....

Pain No. 1

Pain No. 2

☐

☐

☐

☐

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☐

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H7-03 Not including prescribed drugs and other treatments, do you experience pain relief under any conditions like staying still, not wearing a shirt, keeping warm?

☐ No

☐ Yes

Specify the pain alleviating condition(s) .....

.....

Pain No. 1

Pain No. 2

☐

☐

## Non-Painful Sensations

## Section H8

H8-01 Do you suffer from unpleasant sensations that are not painful, e.g. tingling or itch?

☐ No

☐ Yes

Describe the location briefly, e.g. "right hand" .....

and

mark the area affected by those non-painful sensations on page 11. Label them with "S"

H8-02 How would you characterize this sensation?

Specify .....

H8-03 Is the unpleasant sensation always there?

☐ No

☐ Yes

H8-04 If not, how often do you experience this sensation?

☐ Times on average per ☐ day ☐ week ☐ month

Figure 11H

H8-05 Is this non-painful sensation dependent on a stimulus like activity, touch, pressure?

☐ No

☐ Yes

Specify the type of stimulus .....

**Sensory Deficits**

**Section H9**

H9-01 Are there areas where your skin feels numb?

☐ No

☐ Yes

Describe the location briefly, e.g. "right hand" .....

**and**

mark areas of numbness on the drawing on the following page. Label them with "N".

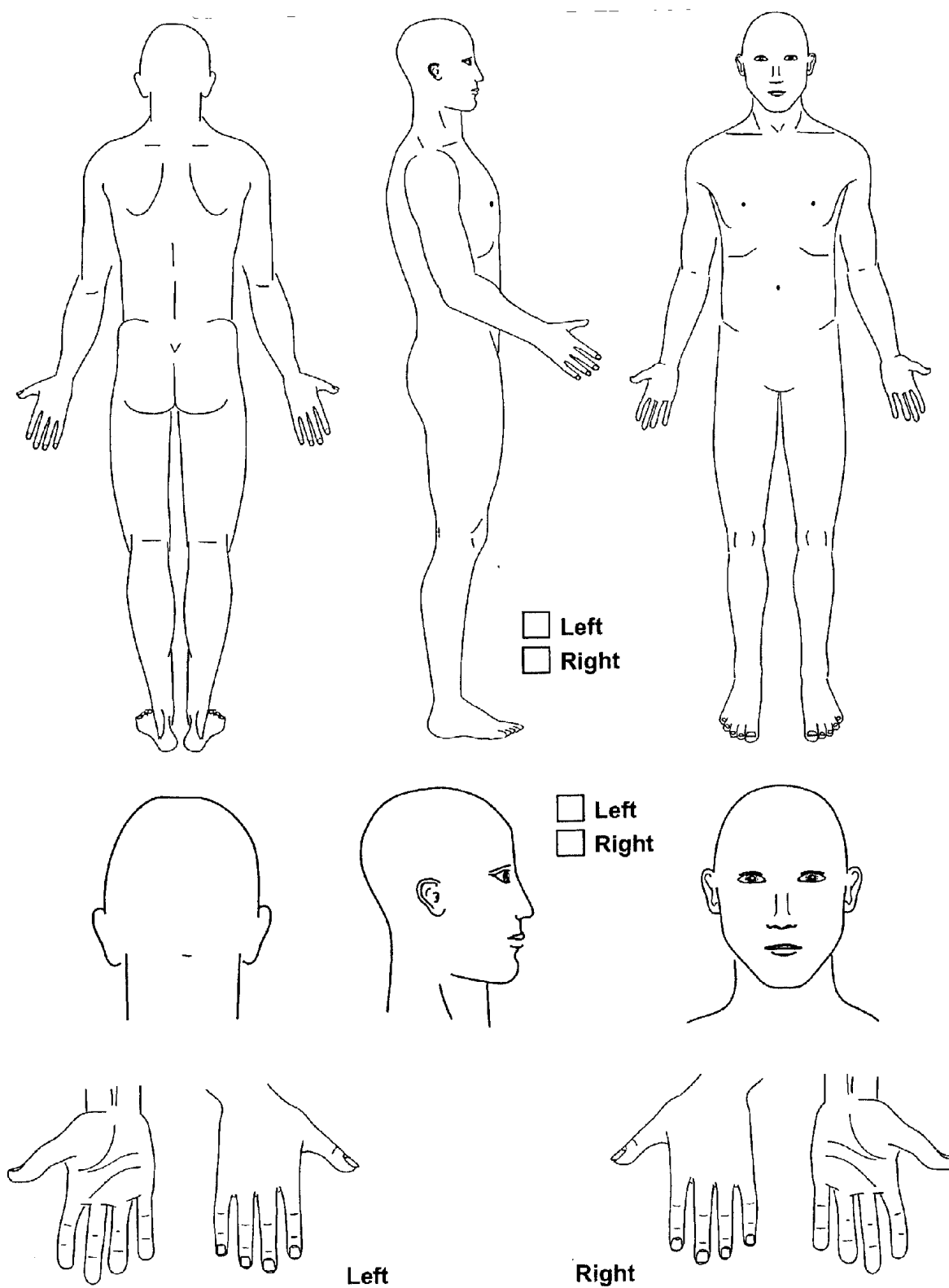


Figure 11J

## ***Clinical Assessment of Pain Syndromes***

### **The Physical Examination**

Date \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_  
(MM/DD/YY)

Patient's First Name \_\_\_\_\_  
Last Name \_\_\_\_\_  
Hospital/Patient Number \_\_\_\_\_  
Date of Birth \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_  
☐ Female      ☐ Male

Investigator's Name \_\_\_\_\_

*Relevant pathological findings made in the general examination by an unblinded physician should be noted separately. A copy of reports concerning special investigations and disease-related findings, e.g. radiological examinations, should be added*

**Skin, Subcutaneous Tissue, and Cutaneous Appendages**

Section S

*The specific location of the findings should be mentioned briefly, e.g. "right hand" or "L4 dermatome". If the findings do not differ in their location, just indicate "same". For some patients it may be appropriate to indicate the distribution of their lesions or other findings in relation to their painful area(s) on the anatomical drawing at the end of the questionnaire*

**S-01 Skin lesions within or related to the painful area**

- ☐ No
- ☐ Yes
- ☐ Wound
  - ☐ Scar
  - ☐ Palpable induration
  - ☐ Neuroma sign (Tinel)
  - ☐ Others
- Specify .....
- Location .....
- Extent, signs of irritation etc. ....

**S-02 Swelling**

- ☐ No
- ☐ Yes
- Location .....

**S-03 Change of skin color**

- ☐ No
- ☐ Yes
- ☐ Reddened
  - ☐ Bluish
  - ☐ Others
- Specify .....
- Location .....

**S-04 Excessive sweating or dry skin**

- ☐ No
- ☐ Yes
- ☐ Excessive sweating
  - ☐ Dry skin
- Location .....

**S-05 Trophic changes of the skin and/or cutaneous appendages**

- ☐ No
- ☐ Yes, involving
- ☐ Skin
  - ☐ Hair
  - ☐ Nails
- Location .....

S-06 **Pressure sensitivity of deep tissues**

*In order to reveal an increased tenderness of subcutaneous, deep tissues within or neighboring the painful area, firm pressure is applied by the examiner's fingertips. The same grip strength has to be tried first on the corresponding area of the non-affected contralateral side to make sure that it is non-painful under normal conditions.*

- ☐ Normal, non-painful sensation of firm pressure  
☐ Increased tenderness or painful sensation elicited

Affected area and tissues (e.g. prominent bones, joints, muscles) .....

.....  
Ask the patient to rate the intensity of the **pressure-evoked pain in deep tissues** by checking one of the four descriptors

- ☐ none      ☐ mild      ☐ moderate      ☐ severe

**and**

on a numerical scale from 0 to 10 ☐

**Nervous System • Sensory System**

Section N

*Sensory stimuli should be applied first to non-affected body parts*

*The specific location of the findings should be mentioned briefly, e.g. "right hand" or "L4 dermatome". If the findings do not differ in their location, just indicate "same". For some patients it may be appropriate to indicate their painful area(s) and, if present, the distribution of their sensory deficits or functional abnormalities on the anatomical drawing at the end of the questionnaire*

**N-01 Sense of touch**

*Two von Frey-filaments, #11 (strength 2.75 g) and #15 (20.9 g) are used. Each filament is applied 4 times. The result is considered positive when 3 out of 4 stimulations produce a response*

☐ Normal (von Frey-filament #11 felt)

☐ Decreased

☐ Mildly (only von Frey-filament #15 felt)

☐ Severely (none felt)

Location .....

☐ Painful sensation elicited by von Frey-filament

☐ #11      ☐ #15

Location .....

Ask the patient to rate the intensity of the **touch-evoked pain** or unpleasant feeling by checking one of the four descriptors

☐ none      ☐ mild      ☐ moderate      ☐ severe

**and**

on a numerical scale from 0 to 10 ☐ where 0 represents no pain and 10 the maximum possible pain

**N-02 Blunt pressure on the skin**

*The blunt eraser end of a pencil is used pressing lightly so as just to indent the skin for 10 seconds. To avoid the sensation of cold, this end must not be made of metal*

☐ Normal sensation

☐ Decreased sensation

☐ Mild decrease (reduced pressure sensation when compared with the non-affected reference area)

☐ Moderate decrease (no pressure, only touch felt)

☐ Severe decrease (no sensation)

Location .....

☐ Painful sensation elicited

Location .....

Ask the patient to rate the intensity of the **blunt touch-evoked pain** or unpleasant feeling by checking one of the four descriptors

☐ none      ☐ mild      ☐ moderate      ☐ severe

**and**

on a numerical scale from 0 to 10 ☐

**N-03 Dynamic stimulus**

*A make-up powder brush (width 1 cm) is lightly moved over the skin at about 3-5 cm per second, in a constant direction*

*Item N-03 continued on the following page*

Item N-03, continued

- ☐ Normal feeling
- ☐ Decreased feeling
- ☐ Mild decrease (reduced movement sensation, but discrimination of the direction preserved)
- ☐ Moderate decrease (vague movement sensation without discrimination of the direction)
- ☐ Severe decrease (no movement felt)

Location .....

- ☐ Painful sensation elicited

Location .....

Ask the patient to rate the intensity of the **brush-evoked pain** or unpleasant feeling by checking one of the four descriptors

- ☐ none      ☐ mild      ☐ moderate      ☐ severe

**and**

on a numerical scale from 0 to 10 ☐

**N-04 Pinprick pain**

*Pinprick is performed with a safety pin*

- ☐ Normal
- ☐ Decreased
- ☐ Mild decrease (less intense pricking sensation when compared with the non-affected reference area)
- ☐ Moderate decrease (no pricking, only blunt touch felt)
- ☐ Severe decrease (no sensation)

Location .....

- ☐ Pinprick evokes a painful sensation that exceeds the intensity of pain felt in the reference region

Location .....

Ask the patient to rate the **increased pinprick intensity** or unpleasant feeling by checking one of the four descriptors

- ☐ none      ☐ mild      ☐ moderate      ☐ severe

**and**

on a numerical scale from 0 to 10 ☐

**N-05 Sense of vibration**

*A standard tuning fork (128 Hz) is used*

- ☐ Normal
- Specify where the sense of vibration was tested .....

- ☐ Decreased
- Specify where the sense of vibration was decreased .....

If using a tuning fork with weights scaled from 1/8 to 8/8, the extent of the decrease should be given by the lowest vibration intensity detected .....

**N-06 Sense of temperature**

*Thermal stimuli can be applied for 10 seconds using either a Peltier-type thermoelectric system set to 20° for the cold stimulus and 40°C for the hot stimulus, or brass bars (diameter 1 cm) can be used. These bars should be stored in thermos vacuum flasks filled with water of the corresponding temperatures*

Item N-06 continued on the following page

Item N-06, continued

- ☐ Normal (warm and cold stimuli discriminated)
- ☐ Decreased
  - ☐ Only for cold
  - ☐ Only for warm
  - ☐ For both, warm and cold

Location .....

- ☐ Unpleasant or painful feeling elicited
  - ☐ Only by cold stimulus
  - ☐ Only by warm stimulus
  - ☐ By both, warm and cold stimuli

Location .....

Ask the patient to rate the intensity of the **temperature-evoked pain** or unpleasant feeling by checking one of the four descriptors

- ☐ none      ☐ mild      ☐ moderate      ☐ severe

and

on a numerical scale from 0 to 10

**N-07 Temporal summation**

*For testing temporal summation, the von Frey-filament #15 (strength 20.9 g) should be used. If this strength elicits pain, the filament #11 (2.75 g) should be employed instead. The filament should be pricked repetitively on the skin at a rate of approximately 3 times a second. Stimulation should last for 30 seconds*

Strength of the von Frey filament used:

- ☐ #11      ☐ #15
- ☐ Sensation remained unchanged
- ☐ Stimulation initially not painful, but pain appeared during stimulation
- ☐ Stimulation already painful at the beginning, but pain increased

**N-08 Sense of position and passive movement**

- ☐ Normal
- ☐ Decreased
  - ☐ Mild decrease (correct discrimination of position and the direction of passive movement is difficult, but still possible)
  - ☐ Moderate decrease (only passively moved body part, but not position or movement direction identified)
  - ☐ Severe decrease (unable to identify passively moved body part)

Specify the limb and the most-proximal joint affected .....

- ☐ Passive movement causes pain

Specify the joint affected ... ..

... and the movement that evokes pain .....

- ☐ Raised leg sign (should be tested only in those patients with low back pain or pain in the lower extremities)

Ask the patient where the movement-evoked pain is felt

- ☐ At the joint
- ☐ At the bones of the limbs moved
- ☐ At tendons or ligaments

Item N-08 continued on the following page

*Item N-08, continued*

- ☐ In muscles
- ☐ At the trunk, e.g. the back or the neck
- ☐ Pain is radiant

Specify .....

Ask the patient to rate the **intensity of pain elicited through passive movement** by checking one of the four descriptors

- ☐ none
- ☐ mild
- ☐ moderate
- ☐ severe

**and**

on a numerical scale from 0 to 10 ☐

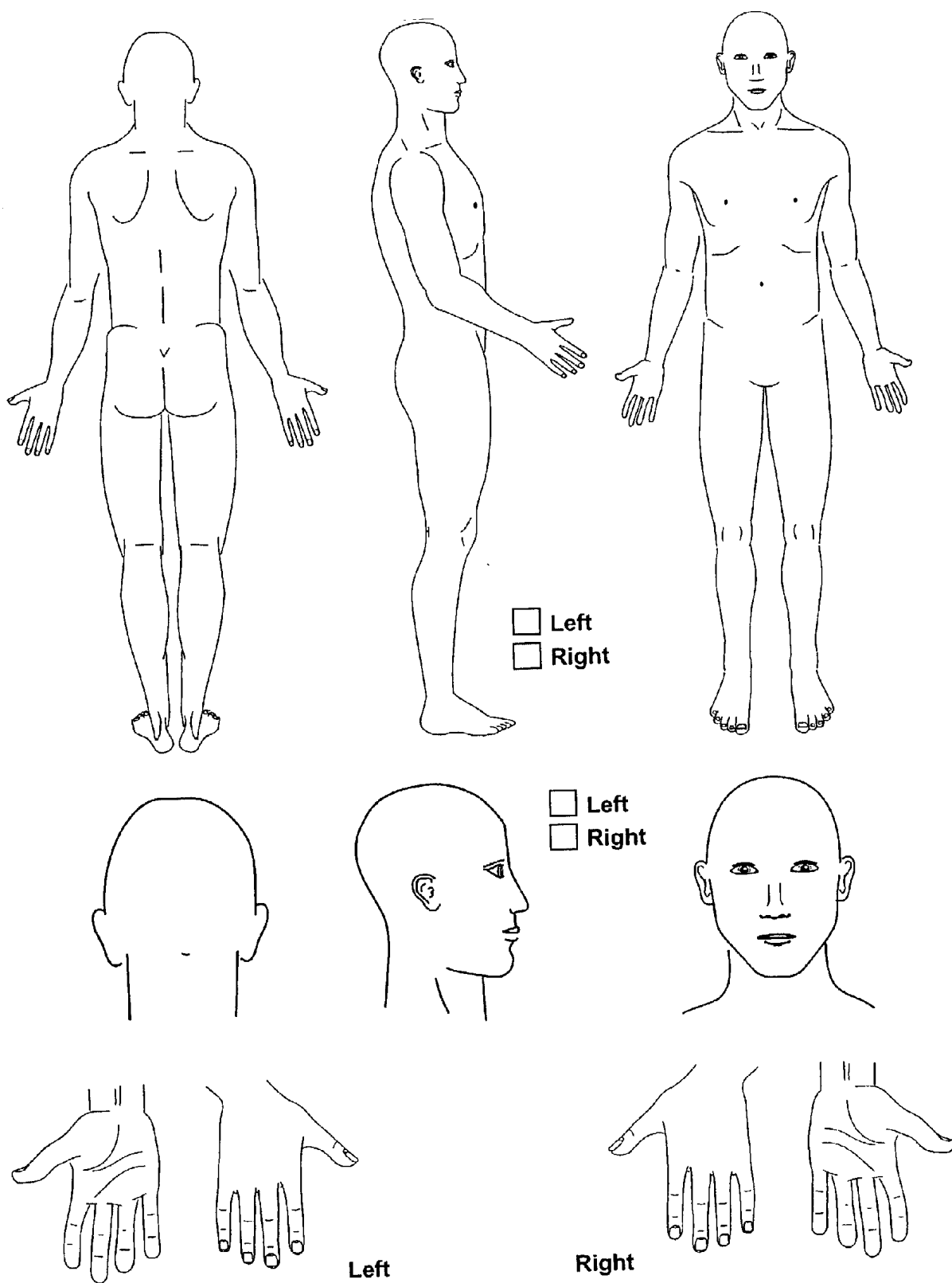


Figure 12H

## PAIN ASSESSMENT

### TECHNICAL FIELD

[0001] This description relates to pain assessment.

### BACKGROUND

[0002] Conventionally, pain syndromes are classified according to the affected anatomical structure, e.g. trigeminal neuralgia, tissue category, e.g. fibromyalgia, or the underlying disease, e.g. cancer pain. They are further divided into acute or chronic conditions related to the duration. Another common division has been to differentiate pain produced by damage to or dysfunction in the nervous system, neuropathic pain, from pain arising from soft tissue, viscera, bone or joints, which have been termed nociceptive or inflammatory pain, depending on the presence or absence of tissue damage and inflammation. There have been attempts to assign specific pain qualities to these different clinical diagnoses. However, the list of qualities attributed to conditions of neuropathic pain alone includes descriptions as diverse as burning, shooting, crawling pain, unusual tingling, electrical sensation or sharp, pulling, aching, tender pain and such descriptors provide little insight into the mechanisms underlying the induction or maintenance of the pain.

[0003] There are several lines of evidence indicating that it is not possible to regard pain in general, or even a particular diagnostic disorder like postherpetic neuralgia, as a uniform entity. The relation between etiology, underlying mechanisms and the specific symptoms and signs related to painful disorders is complex. Pain in an individual patient may be produced by more than one mechanism. No pain mechanism is the inevitable result of a particular etiology. Finally, different mechanisms may operate in patients with the same disease and these may even change during the course of the disease.

[0004] Current strategies for assessing pain severity or the efficacy of a given treatment against pain generally rely on global outcome measures. Patients are asked to rate the overall intensity of their pain on categorical scales ranging from no pain to severe pain. Also widely used are numerical rating scales from 0 to 10, where 10 stands for the worst possible pain, or visual analogue scales, where pain intensity has to be indicated on a horizontal line or similar depiction scaled from 0 to 100. Other, secondary, measures of pain intensity are based on the consumption of analgesic drugs or pain-related deficits in function or the quality of life.

### SUMMARY

[0005] In one aspect, the invention features a method including accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and analyzing the information to determine one or more pain mechanisms indicated by the information.

[0006] Some implementations may include one or more of the following features. The method further includes deriving a pain profile based on the information. The method further includes analyzing the pain profile to determine the presence of at least one pain feature. The pain mechanism is deter-

mined based on the state of the pain feature. The information includes the location of the pain or sensation as indicated on an anatomical diagram. The information includes the depth of the pain relative to the skin surface of the patient. The information pertains to a single pain. The information pertains to two or more separate pains and/or sensations the information being selected to be indicative of one or more than one mechanism associated with the pains and/or sensations. The information includes how long ago the pain or sensation started. The information includes the time course of the pain or sensation, e.g., changes of the pain or sensation over time. The information includes the temporal characteristics of the pain or sensation, e.g., the time profile of instances of the pain or sensation. The information pertains to a sensation other than pain. The information pertains to events that evoke a pain or other sensation, e.g., mechanical or thermal stimuli. The information pertains to the quality of the pain. The information pertains to factors that alleviate a pain or other sensation experienced by the patient. The information pertains to the presence of a sensory deficit. The information is accumulated using a questionnaire and/or by eliciting the patient's history using standardized questions. The information is accumulated by performing a physical examination according to a standardized protocol. The method further includes diagnosing a pain syndrome based on the mechanisms determined. The method further includes treating the patient based on the mechanisms determined. The method further includes analyzing the information along with information accumulated from other patients.

[0007] In another aspect, the invention features a method including accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and graphically or numerically representing a presence, relative amplitude, and absence of individual symptoms and signs to create a pain profile for the patient.

[0008] The invention also features a method including accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and identifying from the information composites of symptoms and signs that represent pain features that are indicative of a pain mechanism.

[0009] In a further aspect, the invention features a method including accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and performing a therapeutic evaluation of a patient based on the information.

[0010] In some implementations, the method also includes, at a later time, accumulating additional information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and, performing a diagnostic or therapeutic evaluation of the patient based on the additional information.

[0011] In yet another aspect, the invention features a method including accumulating information from patients

regarding pains or other sensations experienced by the patients, the information being selected to be indicative of at least one mechanism associated with the pains or sensations, and performing an epidemiological study with respect to the mechanisms associated with the pains or sensations based on the accumulated information.

[0012] The invention also includes a method including accumulating information from patients regarding pains or other sensations experienced by the patients, the information being selected to be indicative of at least one mechanism associated with the pains or sensations, selecting or modifying a drug under development based on the accumulated information, testing the drug on the patients, accumulating additional information from the patients regarding pains or other sensations experienced by the patients, the information being selected to be indicative of at least one mechanism associated with the pains or sensations, and iterating the testing and accumulating until a marketable drug is reached.

[0013] Among other advantages of the invention are one or more of the following. Pathophysiological targets may be identified for rational, non-empirical treatment in individual patients. Pain conditions of a variety of etiologies may be addressed: traumatic, inflammatory and those associated with a structural lesion or malfunction of the nervous system.

[0014] Other features and advantages of the invention will be apparent from the description and drawings, and from the claims.

#### DESCRIPTION OF DRAWINGS

[0015] FIG. 1 is a flow chart comparing a conventional approach to pain management with a mechanism-based approach.

[0016] FIG. 1A is a flow chart illustrating steps of the mechanism-based approach of FIG. 1.

[0017] FIG. 2 is a block diagram illustrating how single symptoms and signs present in conditions associated with spontaneous neuropathic pain are common to multiple clinical diseases and pain syndromes.

[0018] FIG. 3 is a table illustrating the relationship between pain mechanisms and symptoms and signs.

[0019] FIG. 4 demonstrates, by giving two examples, that pain features are composites of pain symptoms and signs.

[0020] FIG. 5 is an illustration of pain profiles produced by examples of different pain mechanisms, showing the pain features typical of these mechanisms.

[0021] FIG. 6 shows an example of a mechanism-based assessment of pain.

[0022] FIG. 6A is a block diagram of a computer system that is suitable for use in mechanism-based pain assessment.

[0023] FIG. 7 is a table providing an overview of examples of treatment strategies based on pain mechanisms.

[0024] FIG. 8 is a flow diagram illustrating an example of steps used in assessing a patient's pain.

[0025] FIGS. 9 and 10 are diagrammatic representations of methods according to two alternative embodiments of the invention.

[0026] FIGS. 11A-11J are pages of an example of a Patient History Questionnaire.

[0027] FIGS. 12A-12H are pages of an example of a Physical Examination Questionnaire.

#### DETAILED DESCRIPTION

[0028] As shown in the two left-hand columns of FIG. 1, conventional management of pain (10) generally relies on clinical diagnoses of diseases and clinical syndromes (16), for example rheumatoid arthritis and herpes zoster, and on knowledge of the underlying etiological or causative factors (11), i.e., in these examples an autoimmune reaction and a viral infection, respectively. The treatment would be aimed at (a) modifying the underlying etiological factors (11) (i.e., disease-modifying therapy (15)), (b) prescribing an analgesic based on empiric standards (i.e., symptom-relief therapy (18)), or (c) a combination of these therapies (12). In this conventional approach, the pain mechanism (13), which actually drives the pain in a patient, would not be identified nor would it be the target of the therapeutic intervention.

[0029] However, as illustrated in FIG. 2 for conditions that are typically associated with spontaneous neuropathic pain, individual symptoms and signs (14) are common to many different clinical diseases or syndromes (16). Thus, neither a disease diagnosis nor individual symptoms alone can be used to identify the mechanisms responsible for the production of the pain.

[0030] As an example, again referring to FIG. 2, the clinical syndrome of nerve entrapment (20) is associated not only with the etiological factor of mechanical compressive injury (22) but also with other etiological factors (11) such as inflammatory changes (26) or the release of cytokines (28) at the site of the injury, or neurodegeneration (30) if untreated, as well as with a possible involvement of the sympathetic nervous system. (Examples of etiological factors (11) are depicted on top of FIG. 2; the lateral extension of the boxes indicates the relevance of each etiological/causative factor for the clinical diseases or syndromes below, as indicated by dotted lines in FIG. 2 for the nerve entrapment example).

[0031] The relationship between clinical diseases and syndromes (16), and spontaneous symptoms and signs (14), is indicated in FIG. 2 by the lines connecting individual diseases/syndromes with the associated symptoms and signs below. Again using nerve entrapment as an example, spontaneous symptoms (14) associated with nerve entrapment include manifestations of spontaneous pain (31), non-painful but unpleasant, disturbing sensations (so-called paresthesia or dysesthesia (32)), ongoing pain (34) or pain that is intermittently present in brief attacks (36). Especially in those cases when the median nerve is entrapped at the wrist, a condition called carpal tunnel syndrome, signs indicating an involvement of the sympathetic nervous system (38) can be found in the physical examination. Altered sweating (40) and a bluish or otherwise changed skin color (42) are indicators of a dysfunction in the sympathetic nervous system, while swelling (44) and trophic changes of the skin or the nails (46) represent consequences of disturbances in the blood supply.

[0032] As illustrated in FIG. 2, symptoms and signs attributable to a dysfunction in the sympathetic nervous

system (38), which can be present in patients with nerve entrapment, as discussed above, can also be present in patients with other diseases, e.g., herpes zoster (58), when the pain that accompanies the acute virus infection turns into a chronic condition named postherpetic neuralgia. Furthermore, symptoms and signs indicative of an involvement of the sympathetic nervous systems may be observed in diseases and injuries that affect structures of the central nervous system, e.g., the brain with stroke (50), the spinal cord after injury (52), and the brain and/or the spinal cord with multiple sclerosis (54). Similarly, the same symptoms and signs involving manifestations of spontaneous pain (31) that are present in patients with nerve entrapment can also be present in patients with other diseases, such as diabetes mellitus (56) and herpes zoster (58).

[0033] Pain mechanisms are those factors operating within the sensory nervous system that are responsible for producing pain symptoms including spontaneous and evoked pain. Different diseases or clinical syndromes may initiate different mechanisms, or the same mechanism, and the same disease or clinical syndrome may generate pain by different mechanisms, either between patients, or even in the same patient over the course of the disease. Thus, patients may have pain mechanisms in common even though their clinical diagnoses are different. For example, a reduced threshold of nociceptive sensory neurons may result from joint inflammation in one patient with, e.g., rheumatoid arthritis, or it may be caused by a reactivation of the varicella-zoster virus in another patient.

[0034] A disease or clinical syndrome (16) (e.g. painful polyneuropathy in diabetes mellitus, or postherpetic neuralgia developing after herpes zoster) is not equivalent, therefore, to a pain syndrome (62, FIG. 1) (e.g., postoperative pain, inflammatory pain, fibromyalgia, central pain, deafferentation pain), which is a particular pattern or constellation of pain mechanisms (13). It is useful for a rational approach to the diagnosis and treatment of pain that pain mechanisms be identified and form the basis for a mechanism-based approach (60) to determining treatment strategies, as shown in the two right-hand columns of FIG. 1.

[0035] In some implementations, a patient's pain mechanisms are identified by first identifying a cluster of multiple, pain-related symptoms and signs, which we call a "pain profile" (64), for the patient. Pain-related symptoms are subjective changes noticed or reported by the patient, e.g. pain provoked by a specific movement like walking or the temporal characteristics of pain attacks. Pain-related signs are objective findings that can be obtained by a clinician examining the patient, e.g. signs of inflammation like red-dening or swelling. The pain profile represents a distinct pattern or "fingerprint" of the patient's pain, displaying all the pain-related symptoms reported by the patient during an interview and the signs elicited by a physical examination.

[0036] Composites of symptoms and signs in the pain profile define "pain features" (66) that each reveal a specific pain mechanism (13) responsible for the development or the maintenance of pain.

[0037] For example, a patient suffering from pain caused by an injury of sensory neurons will complain of numbness in the affected skin area, as sensory neurons are the structures in the peripheral nervous system that are responsible for providing information about stimuli applied to the skin,

such as touch or pinprick. If sensory neurons are lost, the sensation of touch is impaired.

[0038] In the physical examination, sensory loss can be assessed by applying standardized stimuli and recording the patient's response to these stimuli. The composite of the pain characteristics reported by this patient, i.e., the description of numbness, and the sensory loss found upon examination, define a pain feature indicative of pain due to the degeneration of sensory neurons. On the other hand, a patient suffering from pain caused by a reduced threshold of nociceptive sensory neurons, which specifically detect painful stimuli such as pinprick, will complain of an increased sensitivity of the skin, and a physical examination will produce findings that demonstrate an enhanced responsiveness to a pinprick or similar, painful stimuli in the affected area. In this case, the composite of the pain characteristics of this patient defines a pain feature indicative of pain due to a reduced nociceptor threshold.

[0039] Thus, as shown schematically in FIG. 1A, the sequence of steps in providing a mechanism-based pain assessment includes: acquiring patient data (210), generating a pain profile for the patient (212), identifying pain feature(s) (214), and, from the pain feature(s), identifying the corresponding pain mechanism(s) responsible for the patient's pain (216). Based on this analysis, a mechanism-based treatment can be selected (218), and the outcome can be assessed using similar methodology (220).

[0040] FIG. 4 illustrates examples of the composite symptoms and signs that may make up two different, exemplary pain features. Peripheral sensitization, which is characterized by a reduced activation threshold of nociceptors, is indicated by the following symptoms and signs: skin lesions or local signs of inflammation, an intact sensory innervation, and as decisive indicators, an increased painful response (hyperalgesia) to heat stimuli and pinprick as well as a painful sensation felt upon stimulation with normally non-painful stimuli like punctate, blunt or dynamic mechanical stimuli. These symptoms and signs are confined to the site of the skin lesion or inflammation. The complex pain feature of central sensitization results from abnormal sensory excitability of central nervous system neurons, synaptic reorganization and loss of inhibition. As in peripheral sensitization, it is associated with pinprick but not with heat hyperalgesia. Usually non-painful mechanical stimuli again elicit pain, but this is a finding that goes beyond the skin injury site. In addition, temporal summation, which means an increasing sensitivity to repeated stimuli over time for touch and pinprick stimulation may be present.

[0041] Information elicited from the patient is selected to determine the absence or presence of particular symptoms and signs. In addition, the relative amplitudes of individual symptoms and signs are determined by using a rating system such as a categorical scale ranging from none or absent to severe, or a numerical rating scale from 0 to 10. Suitable techniques for eliciting this information will be discussed in detail below. The rating systems are not used only to measure global pain intensity. They are employed to judge the relative contribution of a specific pain symptom or sign to the patient's pain profile. Based on this information, the pain profile enables the determination of the presence or absence of the pain features, and their relative significance for the patient's pain, as will be discussed in further detail below.

[0042] The collection of pain-related data may also include findings from special investigations, e.g., testing beyond the scope of the physical examination, like computed tomography (CT) scans of the spine in patients with low back pain, or the results from a laboratory examination of the cerebrospinal fluid and magnetic resonance imaging (MRI) of the brain and the spinal cord in patients with multiple sclerosis.

[0043] The pain experienced by an individual patient may be caused by more than one pain mechanism (13). For example, a patient's pain profile (64) shown in step 2 of FIG. 6 (discussed in detail below) indicates that the patient's pain is caused by sensory neuron degeneration (85), ectopic activity (87), and synaptic reorganization (82) (step 3, FIG. 6).

[0044] Since multiple mechanisms may co-exist, the pain profile may contain one or more than one pain feature. In a stepwise analysis, the distinctive pattern of symptoms and signs is used first, to identify those pain features, and next, to conclude on the single or multiple pain mechanisms present. This is achieved by comparing the pain profile of the individual patient with a database (e.g., databases 70 and 73, FIG. 6A) that is created using the same mechanism-based tool for the assessment of pain from a large population of patients, including those where single mechanisms can be shown or inferred to be present and associated with particular pain features. As described in detail below, the database 73 includes a collection of pain features associated with single pain mechanisms and provides a key for decoding a patient's pain profile to reveal the most likely or predominant mechanism responsible for his pain, or a complex combination of mechanisms that may co-exist in the patient and constitute a pain syndrome. Matches of the individual patient's pain profile with single mechanism-specific pain features allow detection of the presence of the corresponding pain mechanisms. As a final result, a mechanism-based diagnosis of the individual patient's pain syndrome is obtained.

[0045] In some implementations, a questionnaire on the patient's history (e.g., as shown in FIGS. 11A-11J and discussed below), a physical examination guided by standardized instructions (e.g., as shown in FIGS. 12A-12H), and, where appropriate, special investigations, are used to elicit information from the patient. The questionnaire and the physical examination are used to investigate the presence of spontaneous and evoked pains, temporal and spatial evolution of the symptoms, localization, distribution and nature of evoked pain, and other sensory abnormalities and associated changes. The questionnaire and the standardized instructions for the physical examination are designed to provide a detailed pain profile, which is comprehensive, precise, and reflects the relative contribution of each symptom and sign according to a rating scale.

[0046] This mechanism-based assessment of pain will have consequences for the management of pain. Predominant mechanisms of pain may be identified in a given patient and drugs that are known to target the symptoms generated by these mechanisms can be selected for treatment of the patient. Use of the questionnaire and the standardized examination described below, by enabling the pain profile of the patient to be established, will allow clinicians to standardize investigation and documentation of both initial

findings and findings regarding improvement or worsening of a pain syndrome during follow-up visits. Changes in the mechanisms responsible for the pain over the course of a disease can be followed by the appearance or disappearance of different pain features in the pain profile or relative changes in the rating of correlated symptoms and signs.

[0047] In addition, the development and the evaluation of new analgesic drugs will be facilitated. Mechanism-based pain assessment may improve the design of clinical trials, by allowing the investigation of drug effects on specific mechanisms of pain, rather than attempting to measure overall reduction of pain. Recent evidence from fundamental pharmacological research has revealed previously unknown mechanisms of drug action. These include, for example, the effect of cyclooxygenase (COX)-2 inhibitors on the central nervous system, and the binding of gabapentin to the  $\alpha 2\delta$ -subunit of voltage-gated calcium channels (VGCC), which is up-regulated in animal models of neuropathic pain. Mechanism-based pain assessment will also provide tools to select study populations of patients with pain based on underlying mechanisms. Eventually, mechanism-based assessment of pain may radically change the design of pharmacological trials to study specific effects of drugs on pain mechanisms. Likewise, it will have an important impact on the design of investigations involving the significance of known modes of drug action as well as on alterations in the indication for use of particular analgesics determined in the label or package insert, allowed by regulatory authorities such as the Food and Drug Administration (FDA) or European Agency for the Evaluation of Medicinal Products (EMA).

[0048] FIG. 3 illustrates specific examples of symptoms (left side) and signs (right side) that underlie different mechanisms of neuropathic pain.

[0049] One implementation of mechanism-based pain assessment of pain syndromes is shown schematically in FIGS. 5 and 6. FIG. 6A shows a computer system that can be used to carry out the steps shown in FIG. 6. Initially, a database (70, FIG. 6A) that associates signs and symptoms with pain features is created by investigating the presence and the relative contribution (weighting) of symptoms and signs in a large number of patients with e.g., neuropathic pain. The relationship between single symptoms and signs is examined using statistical methods including correlation analysis, cluster analysis, and analysis of variance. This allows a reduction in the number of symptoms and signs necessary to identify corresponding pain mechanisms. Pain features representing those symptoms and signs, that are attributable to a single mechanism, will be identified by statistical techniques, for example by using an explanatory factor analysis.

[0050] As a result, a series of highly specific pain features (66) (each of which is represented by a single row (72) of boxes (74) in FIG. 5) are established, that can be associated with individual pain mechanisms (13) (listed on the left hand side in FIG. 5). Each of these pain features includes a set of single symptoms and signs (14) (each of which is represented by a box (74) in FIG. 5). The symptoms and signs can be grouped in various ways, such as according to the sequence of their investigation in the questionnaire and in the standardized physical examination. In FIG. 5, the absence or presence of a single symptom or sign is depicted

graphically as an empty box (76) or a filled box (78), respectively. However, absence or presence may be depicted using any suitable graphical indication such as histograms, lines of different thickness, or different symbols. The relevance of an indicative symptom or sign is given, in this example, by a dashed square (80) for a possible indicator and a black square (78) for a strong indicator (this also can be graphically represented in any of a number of different ways). In the database 70, the relevance can be expressed by numbers according to the assigned rating score.

[0051] As discussed above, a patient's pain profile is analyzed to detect the presence of pain features. In a patient whose signs and symptoms are due to the presence of a single pain mechanism, the pain profile is identical to the specific pain feature for that single underlying mechanism. FIG. 5 illustrates pain profiles in hypothetical patients having a single pain feature due to the presence of a single pain mechanism. However, most pain patients will have multiple pain mechanisms operating, and in consequence their pain profile will be a complex composed of multiple pain features whose individual elements may overlap. Here, the database (70) can be used as a "key" to decipher the pain profile "code".

[0052] FIG. 6 illustrates the pain profile (64) of a particular patient with postherpetic neuralgia. Step 3 of FIG. 6 illustrates how, by analyzing the pain profile, three pain features (66), represented by the three rows of boxes in Step 3, can be shown to make up this patient's pain profile. These pain features reflect the concurrence of three different pain mechanisms (13) in this patient that together contribute to the overall pain syndrome (62), i.e., deafferentation pain in this example.

[0053] In Step 1, pain-related symptoms are documented in the patient's history and objective signs are investigated by a physical examination under the standardized conditions of a mechanism-based assessment of pain. In Step 2, the information obtained on the absence or presence, and the rating scores of symptoms and signs is transferred into a graphical (as shown in FIG. 6), or numerical, display representing the patient's pain profile (64).

[0054] In Step 3, the patient's pain profile is analyzed to determine the presence of pain features. Statistical analysis of the individual patient's pain profile can be used to identify single pain features, and in this way allow determination of how many and which mechanisms are responsible for the pain the patient experiences. As outlined above, a database 70 is used to define reference pain features. Matching the individual patient's pain feature with this reference (the reference pain features shown in FIG. 5) produces an estimated factor score. The estimated factor score can be used to assess the individual patient's standing on the corresponding pain feature in the database and, implicitly, on a single underlying pain mechanism. The higher an estimated factor score, the more likely is the presence of the corresponding pain feature and consequently, pain mechanism. The inclusion of a weighting system using categorical, numerical or visual analogue rating scales substantially improves the ability to assess the relative importance or contribution of a single pain mechanism in the patient's pain syndrome and thus, to determine the predominant pain mechanism(s). In FIG. 6, matches are indicated in Step 3 by larger boxes drawn around the small, symptom-indicating

boxes in the top row, and corresponding larger boxes drawn around the matching small boxes in the lower rows. A line is drawn from each of the upper large boxes to a corresponding lower large box to indicate a match.

[0055] In the example shown in FIG. 6, there is a good match of a pain identified by this patient with the pain feature of synaptic reorganization (row 91, FIG. 5), which is characterized by evoked pain. Synaptic reorganization within the central nervous system, due to a sprouting of cutaneous A fibers, is characterized by a location of the pain at the surface (indicated by filled-in box 84 in FIG. 6), rather than in deep tissues. Reorganization means that non-nociceptive sensory information from the periphery, e.g., a touch of the skin, is conducted to relay neurons in the central nervous system that under normal conditions receive information only from nociceptive sensory neurons. As a consequence, touch is now erroneously interpreted as pain (indicated by filled-in box 86 in FIG. 6). Since this mechanism involves structures both of the peripheral and the central nervous system, the pain distribution may follow the territory of a peripheral nerve or it may match a distribution reflecting a central nervous system (CNS) pattern. Therefore, these items are not discriminative for this specific mechanism (empty boxes 221 to 223 in FIG. 6). The affected patients will report pain that is evoked by previously non-painful stimuli like light touch or pressure, and they may also describe limited areas of highly increased sensitivity for evoked pain, so-called trigger zones. In correspondence with the descriptions provided in the patient's history, the physician will find that usually non-painful stimuli like touch, blunt pressure or a (dynamic) stimulus moved over the skin cause pain (indicated by filled-in boxes 86, 88 and 90, respectively).

[0056] Other signs and symptoms exhibited by this patient, e.g., a decrease in the sense of touch, pinprick, vibration and temperature, correspond with the pattern of sensory deficit that is expected to be seen in the presence of sensory neuron degeneration (85), as revealed by comparison with the corresponding pain feature (66) from the database (i.e., the top row (72) of the idealized pain profile shown in FIG. 5).

[0057] Some of the patient's symptoms, e.g., an ongoing pain (box 93) of an aching quality (box 95), referred to deep tissue (box 97), match with the mechanism of ectopic activity. However, this is not exclusively so, as these symptoms are shared by other mechanisms.

[0058] Thus, in this example three mechanisms are present, with sensory neuron degeneration and synaptic reorganization being relatively dominant, and ectopic activity being less relevant. Based on this constellation of pain features, indicating a specific combination of pain mechanisms, the pain syndrome of deafferentation pain is diagnosed in Step 4.

[0059] A patient may have more than one pain and this mechanism-based assessment can be performed for each pain to identify if the same or different mechanisms are responsible.

[0060] The simple absence or presence of individual signs and symptoms may be insufficient to determine what pain mechanisms are responsible for a patient's pain. For example, two patients may both complain about spontane-

ous pain and touch-evoked pain, and one of these patients may also complain of activity-evoked pain. Patient A rates his spontaneous pain as high as 6 on the numerical rating scale (NRS) while touch-evoked pain is only slight and reaches not more than NRS 3. Patient A is more affected by activity-evoked pain (NRS up to 5) than touch-evoked pain. Patient B rates his spontaneous pain NRS 4, but suffers from severe touch-evoked pain of NRS up to 10. Activity-evoked pain is absent. In this example, Patient A's pain is due to the presence of phenotypic changes whereas Patient B's predominant pain is due to central sensitization and synaptic reorganization. Thus, to differentiate between pain mechanisms, symptoms and signs have to be weighted to explore the significance of pain mechanisms in individual patients. This example also demonstrates that reaching a conclusion on the presence of a pain feature may require using the complete information about the absence or presence and the rating of all symptoms and signs examined.

[0061] One application of mechanism-based pain assessment is shown schematically in FIGS. 8 and 9. When a patient (100) visits a clinician (102), e.g., a pain specialist, the clinician, or a nurse or assistant (109), fills in a Patient Data Sheet (104), containing, for example, identifying information (99), previous clinical diagnoses (101), medications (103) and therapies (105) previously prescribed, and relevant special investigations (107). Next, the clinician or a nurse or assistant interviews the patient and completes a standardized Patient's History Questionnaire (106) based on the information provided by the patient. The Patient's History Questionnaire, an example of which is shown in FIGS. 11A-11J and discussed below, is designed to include information on the patient's current pain state (110), the location of the pain (111), onset (112), time-course (113), and temporal characteristics (114) of the pain. Other questions relate to how the pain is evoked (115), the quality of the pain (116), factors that tend to alleviate the pain (117), non-painful sensations (118), and sensory deficits (119).

[0062] After filling out the Patient's History Questionnaire, the clinician conducts a physical examination (108) of the patient, guided by a standardized Physical Examination Questionnaire, which specifies tests that should be performed. The Physical Examination Questionnaire begins with an assessment of changes in the skin, subcutaneous tissue, and cutaneous appendages (120). However, a main focus of The Physical Examination Questionnaire is the investigation of the nervous system and the sensory system (130) in particular. Guided by the Questionnaire, the clinician thoroughly examines, for example, the patient's responses to specific mechanical stimuli (132) including touch, pressure (134), a brush moved over the surface of the skin to elicit dynamic stimulation of nerve fibers (136), and pinprick (138). The Physical Examination Questionnaire is discussed in more detail below, with reference to FIGS. 12A-12H.

[0063] The information obtained by the clinician and the data-taker is stored in a database 71 of patient data and patient profiles (FIG. 6A). The symptoms and signs are used to construct a pain profile for the patient. The pain profile is then analyzed by computer 173 (FIG. 6A) to detect pain features and pain mechanisms, as discussed above. This analysis provides a standardized evaluation of the patient's information (150, FIG. 8). Based on this evaluation, the

physician can arrive at (or computer 173 can generate) a mechanism based strategy for treatment of the patient's pain (152, FIG. 8).

[0064] As described above and illustrated in FIGS. 1 and 8, a mechanism-based assessment of pain enables determination of the pain mechanisms present in the patient and use of these mechanisms as targets for the treatment of pain. So far, few treatment options exist that are aimed at the specific mechanisms underlying a patient's pain. Some existing drugs (180) that can be assigned to particular pain mechanisms (13) are listed in FIG. 7. These include e.g., sodium channel blockers like local anesthetics, carbamazepine, and lamotrigine, which reduce ectopic activity. However, knowledge about the effect of drugs on mechanisms involved in the production or maintenance of pain is derived from animal models. Current clinical assessment of pain does not enable identification of pain mechanisms. For example, carbamazepine or lamotrigine are currently prescribed based on a clinical diagnosis, as for trigeminal neuralgia, or empirically to treat brief pain attacks that are similar in terms of duration and intensity to those pain attacks in trigeminal neuralgia. Or they are used as an additive treatment option for neuropathic pain in general, when pain appears resistant to treatment with other drugs. The majority of drugs listed in FIG. 7 are compounds that have not been used in the treatment of patients but represent substances that have been effective in animal models of pain, like inhibitors of MAPK/ERK in conditions associated with an increased excitability of sensory neurons.

[0065] A mechanism-based assessment of pain will improve the transfer of knowledge about the efficacy of compounds derived from animal models into clinical application as it provides a method to design drug trials based on a specific pain mechanism (FIG. 10). Patients may be enrolled into trials only if they have a particular pain feature that is mediated by a particular target for a drug's action. A mechanism-based assessment of pain is required, therefore, to evaluate the specific efficacy of drugs on individual pain mechanisms in a clinical trial (FIG. 10). This is required in order to investigate and compare the effect of both existing analgesic drugs and new analgesics in FIG. 7. If the patient returns for further treatment and follow up care, the Patient's History and Physical Examination Questionnaires can again be used, the updated information stored in the database 70 of patient data and patient profiles and used to compare changes in pain features in the updated pain profile. The evolution or change in the patient's pain profile can be used to monitor changes in the mechanisms responsible for the pain and to determine the susceptibility of the mechanisms to particular treatment strategies. The information obtained during follow up visits can be used to evaluate whether the treatment is in fact reducing the pain, and thus whether the pain medication or dosage should be changed. Because the information is in a standardized format, initial and follow up data can be readily compared. Moreover, data from one patient can be compared to data from other patients, e.g., patients receiving similar pain medication.

[0066] Referring to FIGS. 6A and 9-10, with suitable precautions taken to protect patient confidentiality and after informed consent has been obtained from the patients involved, patient data stored in the patient database 180 (FIG. 9), or the derived pain profiles or pain features stored in research database 182 (FIG. 9), may be shared with

investigators **200** performing pain-related studies, e.g., studies evaluating the efficacy of pain medications (**186**), and/or with pharmaceutical companies **202** specializing in the development (**188**) of pain medications. This shared data may lead to a beneficial exchange of information between the investigators and/or companies and the clinician, with possible benefit to the patient, for example new medications or treatment protocols.

[**0067**] Information obtained from the clinician (and other clinicians) using the Patient's History Questionnaire and the standardized Physical Examination to construct pain profiles and identify pain features, can also be used by pharmaceutical companies in designing new clinical trials performed on participants to evaluate the efficacy of new pain medications. The results of such trials can be stored in research database **182**, and forwarded to the FDA (**204**) in the hope of obtaining approval for the new medications.

[**0068**] The development of new drugs by pharmaceutical companies (**202**) requires the establishment of biological hypotheses about the mechanisms involved in the pathophysiology of pain (**224**), e.g. the induction of COX-2 in the central nervous system in conditions of inflammatory pain. This hypothesis is tested in animal models (**225**), e.g. by administration of COX-2 inhibitors to the central nervous system. A mechanism-based assessment of pain will not only facilitate formulating biological hypotheses, it will also substantially improve the comparability of results obtained in animal studies with findings in drug trials aimed at proving the concept in patients (phase **2** trials) (**226**). Information on pain mechanisms, transferred to a research database, can be used by the FDA to classify painful clinical conditions according to their underlying mechanisms (**228**), and to define new outcome measures (**229**). Standardized outcome measures represent an important component of guidelines (**230**) established by the FDA for the development of drugs. These guidelines support the design of large-scale drug trials, that are performed to assess the efficacy and the safety of drugs in diverse clinical conditions (phase **3**) (**227**).

[**0069**] Investigators may use the Patient History and Physical Examination Questionnaires in their own studies and clinical trials. The data obtained from the study may be reported to the FDA, e.g., if the research is being conducted to obtain FDA approval, and/or may be reported in a journal article.

[**0070**] Investigators and pharmaceutical companies may also use information obtained from clinicians who use the Questionnaires to develop other, similar questionnaires and evaluation methods containing different sets of pain features or pain profiles suited to their needs.

[**0071**] FIGS. 11A-11J show one example of a Patient's History Questionnaire that may be used to elicit, it from the patient the information on pain-related symptoms discussed above. Many other types of questionnaires can be developed to elicit patient information.

[**0072**] In the Patient's History Questionnaire (FIGS. 11A-11J), an introductory question (Section H1, FIG. 11B) explores the patient's current pain state. Subsequently, the Questionnaire asks for the pain location (Section H2, FIG. 11B), which the patient is asked to indicate on an anatomical drawing (FIG. 11C). Many patients describe the presence of

more than one pain type. Consequently, the Questionnaire asks the patient to assign the different pain characteristics to a predominant "Pain No. 1" or a less disturbing or distressing "Pain No. 2." The two types of pain must be distinct in terms of the type of pain sensation, their location or frequency, or their temporal characteristics. If applicable, patients may describe more than two types of pain on separate sheets.

[**0073**] The following section of the Patient's History Questionnaire (Section H3, FIGS. 11B and 11D) investigate the onset and time-course of the pain syndrome, e.g., when the pain started, what seemed to cause it, and whether the pain has changed over time.

[**0074**] In Section H4 (FIGS. 11D-11E), the Questionnaire explores in detail the temporal characteristics of the pain. Questions include, e.g., whether the pain is always present, whether it ever suddenly, spontaneously worsens, how often the pain or an increase in pain is experienced, how long pain attacks last, and whether the patient experiences a series of pain attacks.

[**0075**] In the next section, Section H5 (FIGS. 11E-11G), the Questionnaire asks the patient questions concerning evoked pain, i.e., what events cause the pain. For example, the Questionnaire tries to lead the patient to distinguish between pain evoked by mechanical stimuli, e.g., pressure from touch or clothing, or thermal stimuli, e.g., cold air or a warm bath. The patient is asked to rate the intensity of each specific symptom in order to learn about the respective significance of the symptoms.

[**0076**] In the version of the Questionnaire shown in FIGS. 11A-11, two different rating systems are used to rate the intensity of pain, i.e., categorical ("none, mild, moderate, or severe") and numerical scales ("rate the intensity on a numerical scale from 0 to 10"). Categorical and numerical scales are currently widely used for global pain ratings. To compare their usefulness in the evaluation of specific pain-related symptoms, both types of scales are included in this version of the Questionnaire. Either or both scales may be used, depending upon which is best accepted by both patients and investigators in a given setting.

[**0077**] In contrast to conventional assessment tools, the Patient History Questionnaire only briefly asks for descriptions of the pain quality (Section H6, FIGS. 11G).

[**0078**] Next, the Questionnaire investigates what drugs, non-drug treatments, and other factors tend to relieve the patient's pain, and what drugs and non-drug treatments have been ineffective (Section H7, FIGS. 11G-11H.) Finally, the Questionnaire investigates sensory disturbances other than pain, e.g., itching, tingling, or paresthesiae, and the presence of numbness (Sections H8 and H9, FIGS. 11H-11I). In Section H9, the patient is also asked to mark any areas of numbness in the anatomical drawing on the following page (FIG. 11J).

[**0079**] Care has been taken to standardize the Patient's History interview. Each item of the Patient's History Questionnaire provides a pre-selected set of answering options. If a patient considers the options inappropriate, the patient is given the opportunity to formulate alternative answers by using her or his own words. Similarly, the Physical Examination Questionnaire (FIGS. 12A-12H) includes a set of standardized questions with pre-selected answering options.

Again, the Physical Examination Questionnaire shown in FIGS. 12A-12H is but one example of the many types of questionnaires that can be used to provide a standardized examination. The Physical Examination Questionnaire provides a mechanism-oriented examination that is suitable to be performed both by pain specialists, e.g., clinical pain specialists and investigators in drug trials, and by physicians who do not specialize in pain therapy. To facilitate use by general physicians, the operational instructions for each item of the examination have been kept as simple as possible without losing information.

[0080] As discussed above, the Physical Examination Questionnaire begins with an assessment of changes in the skin, subcutaneous tissue, and cutaneous appendages (Section S, FIGS. 12B-12C). Skin changes, such as wounds or scars, may be relevant as they can give a hint to the etiology of a given pain condition. Furthermore, diseases like the complex regional pain syndrome (CRPS) are characterized by swelling, enhanced or reduced sweating, altered skin color and sometimes trophic changes of hairs and nails. Section S also investigates whether the patient is experiencing pressure sensitivity of deep tissues (FIG. 12C). The questions in Section S are formulated as "yes/no" questions, with pre-selected choices of symptoms if "yes" is indicated.

[0081] The Physical Examination Questionnaire then examines the nervous system and the sensory system in particular (Section N, FIGS. 12D-12G). Guided by the Questionnaire, the clinician thoroughly examines the patient's responses to specific mechanical stimuli including touch, pressure, a brush moved over the surface of the skin to elicit dynamic stimulation of nerve fibers, and pinprick. Each stimulus is defined in detail to make sure that it will be applied in a standardized fashion. For example, sense of touch is measured by applying two von Frey filaments, #11 (strength 2.75 g) and #15 (20.9 g), four times to the patient's skin and observing the response. Responses are rated using categorical scales (e.g., "normal", "decreased sensation", "painful sensation"). If pain is evoked by a given stimulus, the pain is evaluated in the same manner as in The Patient's History Questionnaire, i.e., by applying both a 4-point categorical scale and a numerical scale ranging from 0 (no pain) to 10 (maximum possible pain).

[0082] During the nervous/sensory system evaluation, the clinician also examines the patient's sense of vibration (FIG. 12E), sense of temperature (FIGS. 12E-12F) and sense of position and passive movement (FIGS. 12F-12G). The clinician tests whether the patient's response to repeated administration of calibrated mechanical stimuli changes over time due to temporal summation (FIG. 12F). Optionally, the area affected by relevant findings in any of the items of the Physical Examination Questionnaire may be indicated on an anatomical drawing (FIG. 12H).

[0083] Referring again to FIG. 6, in some implementations a data storage 900 holds databases that are created and used by applications running on a computer 173. The databases include a database 71 that contains identifying information and pain profiles of individual patients, a database 70 that associates symptoms and signs and clusters of them with pain features, a database 73 that associates pain mechanisms with pain features and groups of pain features, a database 902 that associates therapies with pain mechanisms, and a database 904 that stores measures of outcome

for various pain mechanisms. The software 906 running on computer 173 includes a conventional operating system 908, a database shell 910, and pain data management application modules 912. One module 914 manages the input by a user of symptoms and signs and other information to be stored in the databases or that are otherwise required by the system. A module 916 generates pain profiles from the user data. A module 918 extracts pain features from the pain profiles. A module 920 derives pain mechanisms from the pain features and groups of them. A module 922 provides therapy information based on the derived pain mechanisms.

[0084] Although we have described some examples above, other embodiments are also within the scope of the following claims.

What is claimed is:

1. A method comprising

accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and

analyzing the information to determine one or more pain mechanisms indicated by the information.

2. The method of claim 1 further comprising deriving a pain profile based on the information.

3. The method of claim 2 further comprising analyzing the pain profile to determine the presence of at least one pain feature.

4. The method of claim 3 comprising determining a pain mechanism based on the state of a pain feature.

5. The method of claim 1 wherein the information includes the location of the pain or sensation as indicated on an anatomical diagram.

6. The method of claim 1 in which the information includes the depth of the pain relative to the skin surface of the patient.

7. The method of claim 1 wherein the information pertains to two separate pains and/or sensations the information being selected to be indicative of one or more than one mechanism associated with the pains and/or sensations.

8. The method of claim 1 in which the information includes how long ago the pain or sensation started.

9. The method of claim 1 wherein the information includes the time course of the pain or sensation.

10. The method of claim 9 in which the time course includes changes of the pain or sensation over time.

11. The method of claim 1 where the information includes the temporal characteristics of the pain or sensation.

12. The method of claim 11 in which the temporal characteristics include the time profile of instances of the pain or sensation.

13. The method of claim 1 wherein the information pertains to a sensation other than pain.

14. The method of claim 1 wherein the information pertains to events that evoke a pain or other sensation.

15. The method of claim 14 wherein the events include mechanical stimuli.

16. The method of claim 14 wherein the events include thermal stimuli.

17. The method of claim 1 wherein the information pertains to factors that alleviate a pain or other sensation experienced by the patient.

18. The method of claim 1 in which the information is accumulated using a questionnaire.

19. The method of claim 1 in which the information is accumulated by eliciting the patient's history using standardized questions.

20. The method of claim 1 in which the information is accumulated by performing a physical examination according to a standardized protocol.

21. The method of claim 20 wherein the physical examination includes an assessment of changes in the skin, subcutaneous tissue, and cutaneous appendages.

22. The method of claim 20 wherein the physical examination includes an investigation of the patient's sensory nervous system.

23. The method of claim 22 wherein the investigation includes examining the patient's responses to mechanical stimuli.

24. The method of claim 23 wherein the mechanical stimuli include one or more of the following: touch, pressure, a brush moved over the surface of the skin to elicit dynamic stimulation of nerve fibers, and pinprick.

25. The method of claim 22 wherein the investigation includes examining the patient's sense of temperature.

26. The method of claim 22 wherein the investigation includes examining the patient's sense of vibration.

27. The method of claim 1 further comprising diagnosing a pain syndrome based on the mechanisms determined.

28. The method of claim 1 further comprising treating the patient based on the mechanisms determined.

29. The method of claim 1 further comprising analyzing the information along with information accumulated from other patients.

30. The method of claim 18 in which the questionnaire includes questions having a pre-selected set of possible answers.

31. The method of claim 1 wherein the information pertains to the quality of the pain.

32. The method of claim 1 wherein the information pertains to a sensory deficit.

33. A method comprising

accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and

graphically or numerically representing a presence, relative amplitude, and absence of individual symptoms and signs to create a pain profile for the patient.

34. The method of claim 32 further comprising analyzing the pain profile to identify composites of symptoms and signs that represent pain features.

35. A method comprising

accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and

identifying from the information composites of symptoms and signs that represent pain features that are indicative of a pain mechanism.

36. The method of claim 34 further comprising analyzing the pain features to identify pain mechanisms responsible for the pain.

37. The method of claim 35 further comprising diagnosing a pain syndrome based on the pain mechanisms identified.

38. A method comprising

accumulating information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and

performing a diagnostic or therapeutic evaluation of a patient based on the information.

39. The method of claim 37 also including

at a later time, accumulating additional information from a patient regarding a pain or other sensation experienced by the patient, the information being selected to be indicative of at least one mechanism associated with the pain or sensation, and

performing a diagnostic or therapeutic evaluation of the patient based on the additional information.

40. A method comprising

accumulating information from patients regarding pains or other sensations experienced by the patients, the information being selected to be indicative of at least one mechanism associated with the pains or sensations, and

performing an epidemiological study with respect to the mechanisms associated with the pains or sensations based on the accumulated information.

41. A method comprising

accumulating information from patients regarding pains or other sensations experienced by the patients, the information being selected to be indicative of at least one mechanism associated with the pains or sensations,

selecting or modifying a drug under development based on the accumulated information,

testing the drug on the patients,

accumulating additional information from the patients regarding pains or other sensations experienced by the patients, the information being selected to be indicative of at least one mechanism associated with the pains or sensations, and

iterating the testing and accumulating until a marketable drug is reached.

\* \* \* \* \*

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#### 摘要(译)

提供了评估疼痛的方法和问卷。

