

# Pain Medicine

Schedule of Benefits  
for Professional Fees

## ARTHROCENTESIS/ INJECTIONS

| Code   | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|--------|--|--|-----------------------|--|
| 4332   | Arthrocentesis, aspiration and/ or injection; small joint, bursa or ganglion cyst (e.g. fingers, toes) (I.P.)  | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the "Participating Benefit - No Hospital Admission" applies. This benefit covers both the professional and the facility fee.  |
| 4333   | Arthrocentesis, aspiration and/ or injection; intermediate joint, bursa or ganglion cyst (e.g. temporomandibular acromioclavicular, wrist, elbow or ankle, olecranon bursa) (I.P.)                               | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the "Participating Benefit - No Hospital Admission" applies. This benefit covers both the professional and the facility fee.  |
| 4334   | Arthrocentesis, aspiration and/ or injection; major joint or bursa (e.g. shoulder, hip, knee joint, subacromial bursa) (I.P.)  | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the "Participating Benefit - No Hospital Admission" applies. This benefit covers both the professional and the facility fee.  |
| 5624   | Injection, anaesthetic agent, intercostal nerve, single (I.P.)   | Yes  | No                    |  |
| 5625   | Injection, anaesthetic agent, intercostal nerve, multiple, regional block (I.P.)   | Yes  | No                    |  |
| 174334 | Arthrocentesis, aspiration and/ or injection; major joint or bursa (e.g. shoulder, hip, knee joint, subacromial bursa) (I.P.) - 2 aspirations / injections in same episode                                       | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the "Participating Benefit - No Hospital Admission" applies. This benefit covers both the professional and the facility fee   |
| 174335 | Arthrocentesis, aspiration and/ or injection; major joint or bursa (e.g. shoulder, hip, knee joint, subacromial bursa) (I.P.) - 3 or more aspirations / injections in same episode                               | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the ""Participating Benefit - No Hospital Admission"" applies. This benefit covers both the professional and the facility fee |
| 304332 | Arthrocentesis, aspiration and/ or injection; small joint, bursa or ganglion cyst (e.g. fingers, toes) , including ultrasound guidance (I.P.)  | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the "Participating Benefit - No Hospital Admission" applies. This benefit covers both the professional and the facility fee   |
| 304333 | Arthrocentesis, aspiration and/ or injection; intermediate joint, bursa or ganglion cyst (e.g. temporomandibular acromioclavicular, wrist, elbow or ankle, olecranon bursa) including ultrasound guidance (I.P.) | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the ""Participating Benefit - No Hospital Admission"" applies. This benefit covers both the professional and the facility fee |
| 304334 | Arthrocentesis, aspiration and/ or injection; major joint or bursa (e.g. shoulder, hip, knee joint, subacromial bursa) including ultrasound guidance (I.P.)  | Yes  | No                    | Where this procedure is done in a setting which does not require hospital admission, the ""Participating Benefit - No Hospital Admission"" applies. This benefit covers both the professional and the facility fee |

## EEG

| Code | Description   | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|------|---|--|-----------------------|--|
| 5905 | Video telemetric electroencephalogram (EEG) recordings including full clinical evaluation and placement of sphenoidal electrodes      | No   | No                    | For procedure codes 5905 and 5906 the benefit incorporates all in-patient attendance |
| 5906 | Video telemetric electroencephalogram (EEG) recordings including full clinical evaluation following placement of sub dural electrodes | No   | No                    | For procedure codes 5905 and 5906 the benefit incorporates all in-patient attendance |

## EMG

| Code | Description            | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules |
|------|------------------------|--|-----------------------|---------------|
| 5880 | Electromyography (EMG) | Yes  | No                    |               |

## EMG

| Code | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules |
|------|--|--|-----------------------|---------------|
| 5881 | Electromyography (EMG) study, rectal mucosal sensitivity testing | Yes  | No                    |               |

## EPIDURAL

| Code | Description   | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules |
|------|---|--|-----------------------|---------------|
| 3540 | Epidural injection (I.P.)   | Yes  | No                    |               |
| 3541 | Caudal epidural (I.P.)  | Yes  | No                    |               |
| 3542 | Epidural injection, of anaesthetic substances and/ or therapeutic substances, diagnostic or therapeutic under radiological guidance one or more levels at the same session (I.P.) | Yes  | No                    |               |
| 3545 | Epidural infusion with cannula  | No   | No                    |               |

## IMPLANTABLE PUMPS

| Code | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules   |
|------|--|--|-----------------------|---|
| 5038 | Refilling and maintenance of implantable pump or reservoir including access to pump port (I.P.)  | Yes  | No                    | Benefit for implantation and maintenance of pain pumps, procedure codes 5038 and 5039, applies for one of the following clinical indications:<br>(a) Diffuse cancer pain<br>(b) Failed back surgery<br>(c) Osteoporosis<br>(d) Arachnoiditis<br>(e) Axial somatic pain<br>(f) Painful neuropathies<br>(g) Spinal cord injury<br>(h) Spasticity arising from multiple sclerosis or cerebral palsy. |
| 5039 | Implantation of catheter system and reservoir; tunnelled, intrathecal or epidural catheter for long term medication administration via an external pump or implantable reservoir/ infusion pump (I.P.) | No   | No                    |   |
| 5042 | Removal of subcutaneous implantable pump (does not apply to removal of CVC) (I.P.)   | No   | No                    | Does not apply to removal of CVC  |

## NERVES

| Code | Description   | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules |
|------|---|--|-----------------------|---------------|
| 5586 | Destruction by neurolytic agent (chemodenervation of muscle endplate); muscles enervated by facial nerve (e.g. for blepharospasm, hemifacial spasm) | Yes  | No                    |               |

## NERVES

| Code | Description   | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules |
|------|---|--|-----------------------|---------------|
| 5606 | Implantation of neurostimulator electrodes, Vagus nerve | No   | No                    |               |
| 5610 | Sensory nerve, neurectomy                               | No   | No                    |               |
| 5622 | E.C.T. (each session)                                   | No   | No                    |               |

## NEURO STIMULATORS

| Code | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|------|--|--|-----------------------|--|
| 5043 | Removal of spinal neurostimulator pulse generator or receiver, or neurostimulator electrode percutaneous array (s) or plate/ paddle (s) (I.P.)                                 | No   | No                    |  |
| 5044 | Revision including replacement, when performed, or re-positioning of spinal neurostimulator electrode percutaneous array (s) or plate/ paddle (s); includes fluoroscopy (I.P.) | No   | Yes                   | <p>Benefit for the insertion of spinal cord stimulators will be subject to the following criteria being satisfied:</p> <p>(a) Whether or not low or high frequency spinal cord stimulator is used must be specified on the claim form.</p> <p>(b) Prior approval is sought by a consultant recognised by Irish Life Health and who also has a Diploma in Pain Medicine.</p> <p>(c) The procedure is performed in a hospital that is listed in the Irish Life Health Directory of Hospitals.</p> <p>(d) Benefit will be provided for the trial stage and subsequent implantation for members who satisfy the following criteria:</p> <ul style="list-style-type: none"> <li>(i) An observable pathology concordant with the pain complaint</li> <li>(ii) Further corrective surgical interventions are unlikely to relieve the patient's pain</li> <li>(iii) Non interventional or other conservative therapies have failed</li> <li>(iv) Oral medications are not effective or cause intolerable side effects</li> <li>(v) No untreated chemical dependency exists</li> <li>(vi) Psychological clearance has been obtained through a consultant psychiatrist or clinical psychologist registered with the Psychological Society of Ireland</li> <li>(vii) No contra indications to surgery are present (sepsis, coagulopathy)</li> <li>(viii) Trial screening with the proposed therapy is successful</li> </ul> <p>(e) Benefit will be provided for implantation following a successful trial if the procedure is performed for one of the following clinical reasons:</p> <ul style="list-style-type: none"> <li>(i) Failed back surgery</li> <li>(ii) Complications, including leg pain, from unsuccessful multiple lumbar surgery to repair lower back problems</li> <li>(iii) Reflex sympathetic dystrophy</li> <li>(iv) Arachnoiditis</li> <li>(v) Radiculopathies</li> <li>(vi) Chronic refractory angina</li> <li>(vii) Painful neuropathies</li> <li>(viii) Spinal cord injury</li> </ul> <p>(f) Benefit for a day case hospital stay will be provided for the trial stage.</p> <p>(g) Benefit for a three day stay for the implantation stage will be provided.</p> <p>(h) Benefit will be provided for five days for members who proceed immediately following the trial to implantation during a single hospital admission. Note: the relevant documentation to support the precertification application must be submitted to Irish Life Health in advance of treatment. Maximum once every 7 years, stimulator or modulator or battery replacement performed within that period will not be payable. Only for Irish Life Health approved brands of stimulators.</p> |

## NEURO STIMULATORS

| Code | Description   | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|------|---|--|-----------------------|--|
| 5051 | Replacement of spinal neurostimulator pulse generator or receiver direct or inductive coupling (I.P.) | No   | Yes                   | <p>Benefit for the insertion of spinal cord stimulators will be subject to the following criteria being satisfied:</p> <ul style="list-style-type: none"> <li>(a) Prior approval is sought by a consultant recognised by Irish Life Health and who also has a Diploma in Pain Medicine</li> <li>(b) The procedure is performed in a hospital that is listed in the Irish Life Health Directory of Hospitals</li> <li>(c) Benefit will be provided for the trial stage and subsequent implantation for members who satisfy the following criteria: <ul style="list-style-type: none"> <li>(i) An observable pathology concordant with the pain complaint</li> <li>(ii) Further corrective surgical interventions are unlikely to relieve the patient's pain</li> <li>(iii) Non interventional or other conservative therapies have failed</li> <li>(iv) Oral medications are not effective or cause intolerable side effects</li> <li>(v) No untreated chemical dependency exists</li> <li>(vi) Psychological clearance has been obtained through a consultant psychiatrist or clinical psychologist registered with the Psychological Society of Ireland</li> <li>(vii) No contra indications to surgery are present (sepsis, coagulopathy)</li> <li>(viii) Trial screening with the proposed therapy is successful</li> </ul> </li> <li>(d) Benefit will be provided for implantation following a successful trial if the procedure is performed for one of the following clinical reasons: <ul style="list-style-type: none"> <li>(i) Failed back surgery</li> <li>(ii) Complications, including leg pain, from unsuccessful multiple lumbar surgery to repair lower back problems</li> <li>(iii) Reflex sympathetic dystrophy</li> <li>(iv) Arachnoiditis</li> <li>(v) Radiculopathies</li> <li>(vi) Chronic refractory angina</li> <li>(vii) Painful neuropathies</li> <li>(viii) Spinal cord injury</li> </ul> </li> <li>(e) Benefit for a day case hospital stay will be provided for the trial stage</li> <li>(f) Benefit for a three day stay for the implantation stage will be provided</li> <li>(g) Benefit will be provided for a three day stay for members who proceed immediately following the trial to implantation during a single hospital admission. Note: the relevant documentation to support the precertification application must be submitted to Irish Life Health in advance of treatment. Maximum once every 7 years, stimulator or modulator or battery replacement performed within that period will not be payable. Only for Irish Life Health approved brands of stimulators.</li> </ul> |

## NEURO STIMULATORS

| Code | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|------|--|--|-----------------------|--|
| 5984 | Insertion of spinal cord stimulator - trial stage (I.P.) | No   | Yes                   | <p>Benefit for the insertion of spinal cord stimulators will be subject to the following criteria being satisfied:</p> <ul style="list-style-type: none"> <li>(a) Prior approval is sought by a consultant recognised by Irish Life Health and who also has a Diploma in Pain Medicine</li> <li>(b) The procedure is performed in a hospital that is listed in the Irish Life Health Directory of Hospitals</li> <li>(c) Benefit will be provided for the trial stage and subsequent implantation for members who satisfy the following criteria: <ul style="list-style-type: none"> <li>(i) An observable pathology concordant with the pain complaint</li> <li>(ii) Further corrective surgical interventions are unlikely to relieve the patient's pain</li> <li>(iii) Non interventional or other conservative therapies have failed</li> <li>(iv) Oral medications are not effective or cause intolerable side effects</li> <li>(v) No untreated chemical dependency exists</li> <li>(vi) Psychological clearance has been obtained through a consultant psychiatrist or clinical psychologist registered with the Psychological Society of Ireland</li> <li>(vii) No contra indications to surgery are present (sepsis, coagulopathy)</li> <li>(viii) Trial screening with the proposed therapy is successful</li> </ul> </li> <li>(d) Benefit will be provided for implantation following a successful trial if the procedure is performed for one of the following clinical reasons: <ul style="list-style-type: none"> <li>(i) Failed back surgery</li> <li>(ii) Complications, including leg pain, from unsuccessful multiple lumbar surgery to repair lower back problems</li> <li>(iii) Reflex sympathetic dystrophy</li> <li>(iv) Arachnoiditis</li> <li>(v) Radiculopathies</li> <li>(vi) Chronic refractory angina</li> <li>(vii) Painful neuropathies</li> <li>(viii) Spinal cord injury</li> </ul> </li> <li>(e) Benefit for a day case hospital stay will be provided for the trial stage</li> <li>(f) Benefit for a three day stay for the implantation stage will be provided</li> <li>(g) Benefit will be provided for a three day stay for members who proceed immediately following the trial to implantation during a single hospital admission. Note: the relevant documentation to support the precertification application must be submitted to Irish Life Health in advance of treatment. Maximum once every 7 years, stimulator or modulator or battery replacement performed within that period will not be payable. Only for Irish Life Health approved brands of stimulators.</li> </ul> |

## NEURO STIMULATORS

| Code   | Description   | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|--------|---|--|-----------------------|--|
| 5999   | Insertion of spinal cord stimulator - implantation stage (I.P.) | No   | Yes                   | <p>Benefit for the insertion of spinal cord stimulators will be subject to the following criteria being satisfied:</p> <ul style="list-style-type: none"> <li>(a) Prior approval is sought by a consultant recognised by Irish Life Health and who also has a Diploma in Pain Medicine</li> <li>(b) The procedure is performed in a hospital that is listed in the Irish Life Health Directory of Hospitals</li> <li>(c) Benefit will be provided for the trial stage and subsequent implantation for members who satisfy the following criteria: <ul style="list-style-type: none"> <li>(i) An observable pathology concordant with the pain complaint</li> <li>(ii) Further corrective surgical interventions are unlikely to relieve the patient's pain</li> <li>(iii) Non interventional or other conservative therapies have failed</li> <li>(iv) Oral medications are not effective or cause intolerable side effects</li> <li>(v) No untreated chemical dependency exists</li> <li>(vi) Psychological clearance has been obtained through a consultant psychiatrist or clinical psychologist registered with the Psychological Society of Ireland</li> <li>(vii) No contra indications to surgery are present (sepsis, coagulopathy)</li> <li>(viii) Trial screening with the proposed therapy is successful</li> </ul> </li> <li>(d) Benefit will be provided for implantation following a successful trial if the procedure is performed for one of the following clinical reasons: <ul style="list-style-type: none"> <li>(i) Failed back surgery</li> <li>(ii) Complications, including leg pain, from unsuccessful multiple lumbar surgery to repair lower back problems</li> <li>(iii) Reflex sympathetic dystrophy</li> <li>(iv) Arachnoiditis</li> <li>(v) Radiculopathies</li> <li>(vi) Chronic refractory angina</li> <li>(vii) Painful neuropathies</li> <li>(viii) Spinal cord injury</li> </ul> </li> <li>(e) Benefit for a day case hospital stay will be provided for the trial stage</li> <li>(f) Benefit for a three day stay for the implantation stage will be provided</li> <li>(g) Benefit will be provided for a three day stay for members who proceed immediately following the trial to implantation during a single hospital admission. Note: the relevant documentation to support the precertification application must be submitted to Irish Life Health in advance of treatment. Maximum once every 7 years, stimulator or modulator or battery replacement performed within that period will not be payable. Only for Irish Life Health approved brands of stimulators.</li> </ul> |
| 636052 | Removal of implanted neurostimulator                            | No   | No                    |  |

## NEURO STIMULATORS

| Code   | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|--------|--|--|-----------------------|--|
| 636999 | Combined fee for insertion of spinal cord stimulator - trial and implantation stage on same day (I.P.) | No   | Yes                   | <p>Benefit for the insertion of spinal cord stimulators will be subject to the following criteria being satisfied:</p> <p>(a) Whether or not low or high frequency spinal cord stimulator is used must be specified on the claim form</p> <p>(b) Prior approval is sought by a consultant recognised by Irish Life Health and who also has a Diploma in Pain Medicine</p> <p>(c) The procedure is performed in a hospital that is listed in the Irish Life Health Directory of Hospitals</p> <p>(d) Benefit will be provided for the trial stage and subsequent implantation for members who satisfy the following criteria:</p> <ul style="list-style-type: none"> <li>(i) An observable pathology concordant with the pain complaint</li> <li>(ii) Further corrective surgical interventions are unlikely to relieve the patient's pain</li> <li>(iii) Non interventional or other conservative therapies have failed</li> <li>(iv) Oral medications are not effective or cause intolerable side effects</li> <li>(v) No untreated chemical dependency exists</li> <li>(vi) Psychological clearance has been obtained through a consultant psychiatrist or clinical psychologist registered with the Psychological Society of Ireland</li> <li>(vii) No contra indications to surgery are present (sepsis, coagulopathy)</li> <li>(viii) Trial screening with the proposed therapy is successful</li> </ul> <p>(e) Benefit will be provided for implantation following a successful trial if the procedure is performed for one of the following clinical reasons:</p> <ul style="list-style-type: none"> <li>(i) Failed back surgery</li> <li>(ii) Complications, including leg pain, from unsuccessful multiple lumbar surgery to repair lower back problems</li> <li>(iii) Reflex sympathetic dystrophy</li> <li>(iv) Arachnoiditis</li> <li>(v) Radiculopathies</li> <li>(vi) Chronic refractory angina</li> <li>(vii) Painful neuropathies</li> <li>(viii) Spinal cord injury</li> </ul> <p>(f) Benefit for a day case hospital stay will be provided for the trial stage</p> <p>(g) Benefit for a three day stay for the implantation stage will be provided</p> <p>(h) Benefit will be provided for five days for members who proceed immediately following the trial to implantation during a single hospital admission. Note: the relevant documentation to support the precertification application must be submitted to Irish Life Health in advance of treatment. Maximum once every 7 years, stimulator or modulator or battery replacement performed within that period will not be payable only for Irish Life Health approved brands of stimulators.</p> |

## PAIN BLOCK/ INJECTION

| Code | Description   | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules  |
|------|---|--|-----------------------|--|
| 1220 | Botulinum injector for headaches and migraine   | No   | No                    |  |
| 3543 | Percutaneous lysis of epidural adhesions using solution injection (e.g. hypertonic saline, enzyme) or mechanical means (e.g. catheter) including radiological localisation (includes local anaesthesia and contrast when administered), one or more sessions (I.P.) | No   | No                    | Benefit is limited to 2 treatments per year and only for patients with low back pain in post lumbar surgery syndrome |
| 5575 | Injection of trigeminal ganglion via foramen ovale under image guidance (I.P.)  | No   | No                    | Combined Practitioner Fee - may only be claimed by the anaesthesiologist or the surgeon but not both.                |
| 5580 | Destruction by radiofrequency lesioning of trigeminal ganglion via foramen ovale under x-ray guidance via foramen ovale (I.P.)  | No   | No                    |  |
| 5611 | Transforaminal injection of anaesthetic agent, assessment of response and application of steroid if indicated to medial branch nerve or dorsal root ganglion at one or more levels under image guidance (I.P.)  | No   | No                    |  |
| 5615 | Peripheral nerve block for pain control using nerve stimulator or ultrasound guidance (I.P.)  | Yes  | No                    |  |



## PAIN BLOCK/ INJECTION

| Code | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules |
|------|--|--|-----------------------|---------------|
| 5620 | Sympathetic block, under image guidance (I.P.)                               | Yes  | No                    |               |
| 5621 | Intravenous regional block/ sympathectomy by Bier's technique (I.P.)         | No   | No                    |               |
| 5719 | Chemical sympathectomy, lumbar or coeliac plexus under image guidance (I.P.) | No   | No                    |               |

## PULSED RADIOFREQUENCY

| Code | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules |
|------|--|--|-----------------------|---------------|
| 5612 | Pulsed radiofrequency (PRF) lesioning of medial branch nerve or dorsal root ganglion, one or more levels under image guidance with sensorimotor testing (I.P.) | No   | No                    |               |
| 5614 | Peripheral nerve lesioning including pulsed radiofrequency or electrical stimulation (I.P.)  | Yes  | No                    |               |

## RHIZOTOMY

| Code | Description  | Payable with Private Rooms Technical Benefit | Pre-Approval Required | Payment Rules   |
|------|--|--|-----------------------|---|
| 5616 | Per site - first neurodestructive thermal rhizotomy (temperature > 69°C) under image guidance, with sensory and motor testing, three levels, lumbar, sacral or thoracic (I.P.) | No   | No                    | The following information must be provided on the claim form before benefit can be considered for payment:<br>(a) Details of the level(s) that were treated by rhizotomy i.e. L3 to L5 And/or S1 to S3<br>(b) Confirm the temperature used to perform the procedure<br>(c) Side of the spine – left or right                                  |
| 5617 | Per site - first neurodestructive thermal rhizotomy (temperature > 69°C) under image guidance, with sensory and motor testing, three levels, cervical (I.P.)                   | No   | No                    | The following information must be provided on the claim form before benefit can be considered for payment:<br>(a) Details of the level(s) that were treated by rhizotomy i.e. C3 to C5<br>(b) Confirm the temperature used to perform the procedure<br>(c) Side of the Spine – left or right  |
| 5618 | Subsequent procedure 5616 to the same anatomical site, one or more levels, lumbar, sacral or thoracic - less than 18 months after initial procedure (I.P.)                     | No   | No                    | The following information must be provided on the claim form before benefit can be considered for payment:<br>(a) Date of initial treatment<br>(b) Details of the level(s) that were treated by rhizotomy i.e. L3 to L5 And/or S1 to S3<br>(c) Confirm the temperature used to perform the procedure<br>(d) Side of the spine – left or right |
| 5619 | Subsequent procedure 5617 to the same anatomical site, one or more levels, cervical - less than 18 months after initial procedure (I.P.)                                       | No   | No                    | The following information must be provided on the claim form before benefit can be considered for payment:<br>(a) Date of initial treatment<br>(b) Details of the level(s) that were treated by rhizotomy i.e. C3 to C5<br>(c) Confirm the temperature used to perform the procedure<br>(d) Side of the Spine – left or right                 |