

Colorectal: Faecal Incontinence (Adult) 2021/22



| Indicator Reference Number | Domain | Theme | Measure | Rationale | Name of Indicator / Description | Numerator | Denominator | Period Type | Frequency | Data Source Numerator | Data Source Denominator | Interpretation Guidance | Notes | Reporting Period | | | |
|----------------------------|--|------------------|--------------------------------------|---|---|--|--|-----------------|-----------|-----------------------|-------------------------|-------------------------|--|------------------|-----------------|-----------------|-----------------|
| | | | | | | | | | | | | | | Q1 | Q2 | Q3 | Q4 |
| FI01 | Domain 2: Enhancing quality of life for people with long term | Clinical Process | Pelvic Floor MDT | This is a minimum target level and is in line with recommendations in the service specification | Count of patients discussed at the pelvic floor MDT during the reporting period | The total number of patients discussed at the pelvic floor MDT during the reporting period | N/A | 6 month rolling | Quarterly | Provider submitted | N/A | Higher is better | Service specification states minimum of 50 cases discussed at MDT per year | Jan 21 - Jun 21 | Apr 21 - Sep 21 | Jul 21 - Dec 21 | Oct 21 - Mar 22 |
| FI02 | Domain 2: Enhancing quality of life for people with long term conditions | Clinical Outcome | Anal sphincter repairs | NICE states that the surgical options for faecal incontinence include anal sphincter repair, sacral nerve stimulation or neosphincter formation (Graciloplasty or artificial sphincter.) | Count of anal sphincter repairs undertaken during the reporting period | The total number of anal sphincter repairs undertaken during the reporting period | N/A | 6 month rolling | Quarterly | Provider submitted | N/A | Neutral | Elective only Exclude maternal perineal tears First time repair only OPCS - H57.1 (Placement of artificial anal sphincter NEC), H50.1 (Posterior repair of anal sphincter), H50.2 Anterior repair of anal sphincter) - primary procedure only. DIAG - R15X (Primary or secondary) | Jan 21 - Jun 21 | Apr 21 - Sep 21 | Jul 21 - Dec 21 | Oct 21 - Mar 22 |
| FI03 | Domain 2: Enhancing quality of life for people with long term conditions | Clinical Outcome | Sacral Nerve Stimulation - permanent | NICE states that the surgical options for faecal incontinence include anal sphincter repair, sacral nerve stimulation or neosphincter formation (Graciloplasty or artificial sphincter). | Count of patients who receive a permanent Sacral Nerve Stimulation (SNS) for faecal incontinence | The total number of patients who receive a permanent SNS for faecal incontinence during the reporting period | N/A | 6 month rolling | Quarterly | Provider submitted | N/A | Higher is better | Elective only First procedure only OPCS - A70.1 (Implantation of neurostimulator into peripheral nerve) followed by Z11.2 (Sacral nerve) OR A70.4 (Insertion of neurostimulator electrodes into peripheral nerve) followed by Z11.2 (Sacral nerve). DIAG - R15X (Primary or secondary) | Jan 21 - Jun 21 | Apr 21 - Sep 21 | Jul 21 - Dec 21 | Oct 21 - Mar 22 |
| FI04 | Domain 2: Enhancing quality of life for people with long term conditions | Clinical Outcome | Sacral Nerve Stimulation - temporary | Patients who are unsuitable for anal sphincter repair should be given a trial of temporary sacral nerve stimulation. If successful, then permanent sacral nerve stimulation is indicated. If a trial is unsuccessful then the patient can be considered for neosphincter. | Count of patients who receive a temporary Sacral Nerve Stimulation (SNS) for faecal incontinence | The number of patients who receive a temporary SNS for faecal incontinence during the reporting period | N/A | 6 month rolling | Quarterly | Provider submitted | N/A | Neutral | Elective only First procedure only A70.1 (Implantation of neurostimulator into peripheral nerve), A70.4 (Insertion of neurostimulator electrodes into peripheral nerve), Z11.2 (Sacral nerve), Y70.5 (Temporary operations) DIAG - R15X (Primary or secondary) | Jan 21 - Jun 21 | Apr 21 - Sep 21 | Jul 21 - Dec 21 | Oct 21 - Mar 22 |
| FI05 | Domain 2: Enhancing quality of life for people with long term conditions | Clinical Outcome | Sacral Nerve Stimulation | Patients who are unsuitable for anal sphincter repair should be given a trial of temporary sacral nerve stimulation. If successful, then permanent sacral nerve stimulation is indicated. A successful trial is determined by percutaneous nerve evaluation. | Proportion of patients who go on to have permanent Sacral Nerve Stimulation (SNS) implants following a temporary SNS test | Of those patients in the denominator, the number who subsequently have permanent SNS implants | The number of patients receiving a temporary SNS test in the reporting period | 6 month rolling | Quarterly | Provider submitted | Provider submitted | Higher is better | Numerator: Include all patients from denominator cohort where: OPCS - A70.1 (Implantation of neurostimulator into peripheral nerve) followed by Z11.2 (Sacral nerve) OR A70.4 (Insertion of neurostimulator electrodes into peripheral nerve) followed by Z11.2 (Sacral nerve) Denominator: OPCS - A70.1 (Implantation of neurostimulator into peripheral nerve), A70.4 (Insertion of neurostimulator electrodes into peripheral nerve), Y70.5 (Temporary operations), Z11.2 (Sacral nerve) DIAG - R15X (Primary or secondary) | Jan 21 - Jun 21 | Apr 21 - Sep 21 | Jul 21 - Dec 21 | Oct 21 - Mar 22 |
| FI06 | Domain 2: Enhancing quality of life for people with long term conditions | Clinical Outcome | Infection rates | The proportion of patients presenting with infection requiring device removal or surgical revision is one indicator of the quality of the faecal incontinence surgical service | Proportion of patients admitted with an infection requiring device removal or surgical revision within 180 days following permanent Sacral Nerve Stimulation (SNS) implantation, replacement, or revision | Of those patients in the denominator, the number who were subsequently admitted with an infection requiring device removal or surgical revision within 180 days of initial procedure | The number of patients receiving permanent, replacement or revision of SNS in the reporting period | 6 month rolling | Quarterly | Provider submitted | Provider submitted | Lower is better | Numerator: Include all patients from denominator cohort where: OPCS - A70.1 (Implantation of neurostimulator into peripheral nerve), Z11.2 (Sacral nerve) OR A70.2 (Maintenance of neurostimulator in peripheral nerve), Y03.2 (Renewal of prosthesis in organ NOC), Z11.2 (Sacral nerve) OR A70.2 (Maintenance of neurostimulator in peripheral nerve), (Y03.1 (Maintenance of prosthesis in organ NOC), Y03.3 (Correction of displacement of prosthesis NOC), Y03.4 (Other resting of prosthesis in organ NOC) OR Y03.5 (Conversion to prosthesis in organ NOC) OR Y03.6 (Adjustment to prosthesis in organ NOC) OR Y03.8 (Other specified attention to prosthesis in organ NOC) OR Y03.9 (Unspecified attention to prosthesis in organ NOC) Denominator: OPCS - H50.1 Posterior repair of anal sphincter H50.2 Anterior repair of anal sphincter DIAG - T85.7 (Primary) DIAG - R15X (Primary or secondary) | Jul 20 - Dec 20 | Oct 20 - Mar 21 | Jan 21 - Jun 21 | Apr 21 - Sep 21 |

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| | | | | | | | | | | | | | | Q1 | Q2 | Q3 | Q4 |
| FI07 | Domain 2: Enhancing quality of life for people with long term conditions | Clinical Outcome | Bulking Agents | NICE guidance states that the current evidence on the safety and efficacy of injectable bulking agents for faecal incontinence does not appear adequate for the procedure to be used without special arrangements for consent and for audit or research. This should take place in the context of a clinical trial or formal audit protocol that includes information on well-defined patient groups. | Count of patients receiving injecting of bulking agents | The number of patients receiving injecting of bulking agents in the reporting period | N/A | 6 month rolling | Quarterly | Provider submitted | N/A | Neutral | OPCS - H57.8 (Other specified other operations on the anal sphincter to control continence), Y39.3 (Injection of inert substance into organ NOC) DIAG - R15X (Primary or secondary) | Jan 21 - Jun 21 | Apr 21 - Sep 21 | Jul 21 - Dec 21 | Oct 21 - Mar 22 |
| FI08 | Domain 2: Enhancing quality of life for people with long term conditions | Clinical Process | Bulking Agents | NICE guidance states that the current evidence on the safety and efficacy of injectable bulking agents for faecal incontinence does not appear adequate for the procedure to be used without special arrangements for consent and for audit or research. This should take place in the context of a clinical trial or formal audit protocol that includes information on well-defined patient groups. | Proportion of injecting of bulking agent procedures performed as a day case | Of those procedures in the denominator, the number that were carried out as a day case | The number of injecting of bulking agent procedures carried out in the reporting period | 6 month rolling | Quarterly | Provider submitted | Provider submitted | Higher is better | Numerator: Include all patients from denominator cohort where: PATIENT CLASSIFICATION = '2' Denominator: OPCS - H57.8 (Other specified other operations on the anal sphincter to control continence), Y39.3 (Injection of inert substance into organ NOC) DIAG - R15X (Primary or secondary) | Jan 21 - Jun 20 | Apr 21 - Sep 21 | Jul 21 - Dec 21 | Oct 21 - Mar 22 |
| Data collection has been approved by the Review of Central Returns - ROCR ROCR/OR/2230/001MAND | | | | | | | | | | | | | | | | | |