Control your risks, control your outcomes

Is your revision arthroplasty patient at risk of developing a surgical site complication?



Smith-Nephew

PICO^{\$} 7 Single Use Negative Pressure Wound Therapy System

Helping you get **CLOSER TO ZERO**^o surgical site complications¹

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PICO[®] ORTHOPAEDIC RISK QUESTIONNAIRE REVISION

Is your patient high risk?

Operative time

Revision hip arthroplasty (rTHA) can take, on average, 78 mins longer compared with primary procedures²

Revisio

Surgical site infection (SSI) risk can double with rTHA compared with primary procedures³ Deep or organ space SSI ca nearly quadruple with rTHA compared with primary procedures³

SSI risk can double

with rTHA compared with primary procedures³

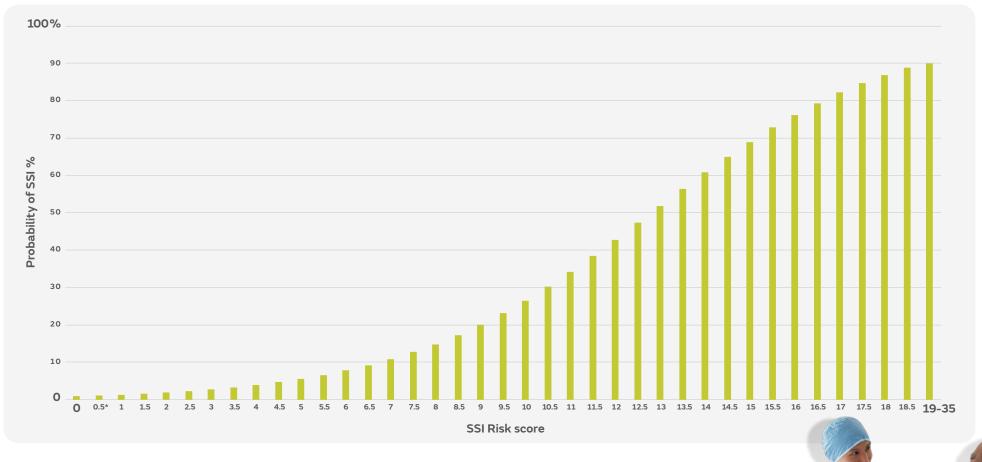
REFERENCE

Is your patient high risk?

Certain patient factors correlate with SSI development following primary and revision arthroplasty⁴. Pre-operative identification can determine the probability of an SSI developing post-operatively⁴.

| | | | ary knee Revis | | Revision knee | Presence | | | |
|--|-----|---|----------------|---|---------------|-------------------|---------------------------------------|--------------------|--|
| Score | 0 | 1 | 3 | | 3 | Score | 1 | 0 | |
| Diabetes | | | Long term in | sulin use | | Rheumatoid ar | thritis or inflan | nmatory arthropath | |
| | | | Presence | | | Presence | | | |
| Score | | | Score | | 0 | Score | | | |
| | | | | | | | | | |
| Tobacco use | | | Lower-extrem | Lower-extremity osteomyelitis or pyogenic arthritis | | | Pelvis, thigh, leg traumatic fracture | | |
| | | | Presence | | | Presence | | | |
| Score | 1.5 | 0 | Score | 2 | 0 | Score | 2 | 0 | |
| Lower-extremity pathologic fracture | | | Morbid obesi | Morbid obesity (BMI ≥ 40) | | | Primary bone cancer | | |
| | | | Presence | | | Presence | | | |
| Score | 2.5 | 0 | Score | 2.5 | 0 | Score | 4 | 0 | |
| Reaction to prosthesis or implant within 3 years | | | Staphylococ | Staphylococcal septicemia | | | TOTAL PATIENT SCORE: | | |
| | | | Presence | | | | | | |
| Score | | | Score | | 0 | | | | |

SSI risk score and corresponding probability of SSI



Certain patient factors correlate with SSI development following primary and revision arthroplasty⁴. Pre-operative identification can determine the probability of an SSI developing post-operatively⁴.

*Interpolated value. A score of 0.5 is not a possible result of any combination of positive risk factors.

CONTROL YOUR RISKS, CONTROL YOUR OUTCOMES | 4

Control your risks, control your outcomes

The prophylactic use of incisional NPWT significantly reduced **LoS** by an average of **1.87 days**^{*} following revision hip and knee arthroplasties²

The risk of developing a post-operative SSC depends on the type of surgery and patient risk factors^{6,7}

The presence of just **1 major risk factor or 2** or more moderate risk factors, places patients at high risk of an SSC and means you should consider **PICO**^o sNPWT⁶



| Category | Patient-related risk factor | Procedural-related risk factor |
|--|---|---|
| | BMI ≥ 40kg/m ² or ≤ 18kg/m ² | Extended surgery [†] |
| | Uncontrolled insulin dependent diabetes mellitus | Emergency surgery |
| Major risk factor presence of 1 = high risk of surgical site complication | Renal dialysis | Hypothermia |
| | I ASA physical status >II | Inaemia / blood transfusion |
| (仏『凵) | Age < 1 year or > 75 years | I High wound tension after closure |
| Moderate risk factor | BMI 30-39.9kg/m ² | Dual antiplatelet treatment |
| presence of 2 ≥ high risk of surgical site complication | Immunosuppression | Suboptimal timing or omission of prophylactic antibiotics |
| | Smoking (current) | Tissue trauma / large area of dissection / large area of undermining |

Table adapted from World Union of Wound Healing societies Consensus, 2016. The risk factors represented in this table are examples only and not an exhaustive list.¹⁴ *Compared with standard care; 6.71 days v 8.58 days; p = 0.019. [†]Defined as >T (hours) which is dependent on the type of surgical procedure, and is the 75th centile of duration of surgery for a particular procedure, e.g. coronary artery bypass graft has a T of 5 hours and caesarean section has a T of 1 hour.

Revision surgery

Dr Thomas Goetzmann, Clinique Louis Pasteur, Essey les Nancy (France) PICO° sNPWT on a surgical incision after rTKA surgery

Patient

A 73 year-old female, the patient was unable to walk unaided.

Class 1 obesity

History of rTKA surgery due to infection; the patient had already received two rTKAs.

The patient underwent a left rTKA and put on antibiotics

PICO sNPWT treatment methodology

- It was decided to apply the PICO sNPWT system in the operating theatre. The chosen PICO dressing size was 10 x 40cm and the incision line measured 21cm L, with sutures present
- The surgeon decided to leave the PICO system in place for 6 days

Course of PICO sNPWT

- The patient was in hospital for 6 days before transfer to post-acute care and rehabilitation.
- The patient's leg was not immobilised and she was able to wear her compression stocking, the PICO system is compatible for use with compression therapy
- At D6, 20% saturation of the dressing was observed
- There was no sign of inflammation under the dressing. After a team discussion, a new PICO dressing (15 x 30 cm) was applied in order to avoid any risk of dehiscence in areas where the peri-wound skin was more fragile, which was distinguishable by a whitish discolouration. The surgeon decided to leave the PICO system in place for another 6 days

CONTRACTOR

The patient was able to receive physiotherapy throughout treatment with **PICO** sNPWT in order to regain the ability to walk unaided

Treatment outcome

- The patient was seen again at D12. The PICO dressing showed 45% saturation, and the sutures at the ends of the incision line were removed. There was no sign of inflammation or dehiscence. The surgeon continued PICO therapy for another 7 days, with the same dressing size
- PICO sNPWT was discontinued at D19. The dressing showed two stains, and all remaining sutures were removed. Antibiotic therapy was discontinued on the same day
- The total duration of treatment was 19 days. The PICO Dressing was then replaced by a hydrocellular dressing
- By D47, the incision was fully healed without any complications
- The patient was pleased with the outcome and with the device, which helped prevent further complications and allowed her to resume physiotherapy sessions in order to regain mobility
- The surgeon was also pleased with the device

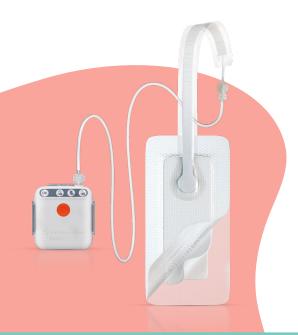
Revision surgery(continued)

Dr Thomas Goetzmann, Clinique Louis Pasteur, Essey les Nancy (France) **PICO**° sNPWT on a surgical incision after rTKA



Evolution of closed surgical incision with PICO^o sNPWT









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For detailed product information, including indications for use, contraindications, precautions and warnings, please consult the product's applicable Instructions for Use (IFU) prior to use.

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