One year outcome of unicompartmental interpositonal arthroplasty of the medial compartment of the knee, utilizing a metallic implant.

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Outline

- Introduction of a metallic implant for interpositional medial compartmental arthroplasty of the knee (UIA)
- Presentation of early results of initial 30 procedures, as a single surgeon case series
- Considerations re. further direction

Introduction

- Medial compartment osteoarthrosis of the knee
- Multiple treatment options
- Escalating commitment :
 - morbidity vs adequacy
 - 'when to burn which bridge'

Introduction

Non-established treatment options

Clinical outcome not known

- Range of possible outcomes
- Decision analysis (COA 2007) can help with preliminary assessment of the balance of risks and benefits
- Initial case series can provide preliminary data re. morbidity and efficacy

Metallic interpositional arthroplasty

Historical:

- → MacIntosh, McKeever
- \rightarrow Sbarbaro, Swanson
- used initially in OA and RA
- as far back as late 1950's
- required some bone preparation

Metallic interpositional arthroplasty

- Unispacer (Sulzer, Zimmer)
- brief appearance in early 2000's
- approximately 3000 implanted
- minimal reporting
- issues: implant instability, pain, stiffness
- relied on femoral congruency for stability
- no longer used

Metallic interpositional arthroplasty

- Dr. R. Scott, Boston \rightarrow McKeever
 - 'may be considered as a bridging measure in the treatment of unicompartmental OA'
 - 70-86% implant survival at 8 y \rightarrow not unlike HTO
 - 10 out of 24 doing well at 16 years (2006)

- Development history
 - 2003 trial of a polyurethane interpositional arthroplasty implant (Advanced BioSurfaces)
- Technique established
- Initial recovery OK
- Implant stable
- Synovitis due to wear after 4-6 months
- Trial stopped

- Development history:
- lessons learned
- metallic implants (\rightarrow 510k)
- 3 and 4 mm implants, various AP sizes
- 2007 (Arnold)
 - 300 implants
 - 92/100 patients with functional outcome scores
 - functional outcome scores encouraging
 - 10% revision rate at 1 y
 - 1/300 dislocation, 1/300 infection.
- to date: approximately 600 implants placed
- insufficient data capture

- Considerations:
- Is it safe?
- Is it effective?
- What about long-term management?
- Is it acceptable to the health care system?
- Cost and other resource utilization?
- Health Canada licencing status?

- Medial implant licenced by HPB, Health Canada in 2009
- Note: changed to Special Access in 2011 (insufficient data)

- Safety:
- less invasive surgery
- minimal hospital stay
- no violation of subchondral bone → potentially 'reversible' (management of infection etc)

OrthoGlide - medial



- Process of patient consent critical
- Patient tolerance of uncertainty of effectiveness of implant vs. assessment of exposure to surgical risks associated with various options
- INFORMED CONSENT of high quality

Female, 80y. Medial UKA 4 years earlier



2 years post-op



2 years post-op



 Period under observation: July 15, 2009 - July 15, 2010

• Thirty patients: 23 men / 7 women

• Average age 63.9 years, range 44 to 87 y

Local anesthesia with IV sedation

 Arthroscopic assessment and partial joint preparation

Arthrotomy (5-7 cm) for completion of joint preparation and implant insertion

• Daycare surgery

• Surgeon follow-up:

-2w, 2m, 6m, 1y and as needed

• Chart review up to and incl 1 year mark

- Outcome at 1 year (surgeon rating):
- 'good' (satisfactory, steady-state, likelihood of early revision low)
- 'fair' (implant merely tolerated, not a steady-state, likelihood of early revision significant)
- 'poor' (implant not tolerated, revision pending or performed).

- All completed as intended
- No conversion to general anesthesia
- No overnight stay or early re-admission.

 One patient lost to follow-up immediately after surgery

• One hemarthrosis: wash-out POD 37

No dislocation

No revision surgery

Knee range of motion

2 months $0-125 (\pm 10)$ degrees

6 months $0-128 (\pm 7)$ degrees

12 months $0-131 (\pm 7)$ degrees

Functional rating at one year.22/29 patients (76%) \rightarrow good3/29 patients (10%) \rightarrow fair4/29 patients (14%) \rightarrow poor

(with patient lost to follow-up assigned to the 'poor' group \rightarrow 73% 'good', 10% 'fair' and 17% 'poor'.)

'Poor' group:

$2/4 \rightarrow$ progression of OA in lateral compartment

$2/4 \rightarrow$ unrelenting discomfort

Offered conversion to TKR

Discussion

After UIA \rightarrow 'good' results at 1 y in 70-80% range (???)

After TKR \rightarrow 'good' results at 1 y in 85-90% range (NIH, CIHI)

After UKR \rightarrow similar or slightly less than TKR?

Discussion

- 'Good' TKA
- VS
- 'Good' UKA
- VS
- 'Good' UIA

Comparative outcome assessment needed -Matched cohort study vs randomized trial???

Conclusion

- Interpositional arthroplasty of the medial compartment of the knee with the metallic Orthoglide implant appears to be safe and can be effective
- Uncertainty persists re. consistency and extent of functional improvement
- Revision options are preserved

Conclusion

- Further assessment will require a structured roll-out with systematic data capture:
 - -on-line data registry with ongoing analysis vs
 - -formal multi-centre trial

Conclusion

 Refinement in implant design and materials, technique, indications etc to be based on further data collection

 Open communication in the orthopaedic community will be required to assess relative merit of various established and emerging technologies

THANK YOU