A descriptive clinical evaluation of arthroscopic synovectomy in rheumatoid knees: a prospective study

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(Index words: Function, pain, perioperative morbidity, range of motion)

Abstract

Objective To evaluate the clinical outcome of arthroscopic synovectomy for persistent rheumatoid synovitis of the knee joints.

Design Prospective clinical study.

Setting Kasturba Medical College Hospital, Manipal.

Patients Fifty two knee joints in 46 patients.

Measurements The effect of the procedure and its influence in the progression of the disease process on knee joints were assessed in terms of reduction of pain, improvement in range of motion, improvement in functional activity and recurrence of synovitis with effusion.

Results During the average follow up period of 5 years, the patients showed appreciable improvement (90% of knee joints) until 3 years of follow up. At the end of 5 years of follow up, it reduced to about 75%.

Conclusions Arthroscopic synovectomy along with medical treatment can control the disease process and preserve the knee joint function for up to 3 years.

Introduction

Descriptions of the surgical excision of the synovium from a variety of joints have been published for over a century. Open synovectomy of the knee, used first for the treatment of tubercular arthritis, was later used in rheumatoid arthritis. The main disadvantage of open synovectomy of the knee is related to the rehabilitation of the patient after the surgical procedure. Most authors recommended a need for manipulation of the knee under anaesthesia during the early post-operative period to overcome loss of motion. Extensive physical therapy and prolonged hospital stay are often necessary to obtain a good result [1]. Wound separation, haematoma, infection and cutaneous nerve injury represent other documented complications.

The enlargement of the field of application of arthroscopy from diagnostic use to therapeutic purposes and the development of new instruments such as the motorised intra-articular shaver have facilitated safe and efficient arthroscopic synovectomy. The main change is in the technique rather than the concept or indications. Our study was designed to clinically evaluate the efficacy, tolerance, and morbidity of arthroscopic synovectomy in a group of patients with persistent active rheumatoid synovitis, by comparison of pre-operative and post-operative pain, range of movement and functional activity, and recurrent synovitis with effusion.

Materials and methods

This is a prospective study of 46 patients (52 knees) of arthroscopic synovectomy of the knee done in Kasturba Medical College Hospital, Manipal, India, from December 1997 to January 2000. Rheumatoid arthritis patients with persistent synovitis in the joint causing pain and swelling despite 6 months of adequate medical management were included in the study. All the patients had loss of functional activity and a reasonable range of motion. Patients with marked destruction and deformity of the joint, and those who had previously undergone open synovectomy were excluded from the study.

After clinical examination and relevant investigations, the diagnosis of rheumatoid arthritis was established. The patients were scored pre-operatively using the criteria of pain, range of motion, functional activity and synovitis with effusion. All operations were done under general or spinal anaesthesia with a pneumatic tourniquet. Synovectomy was performed in both anterior and posterior compartments using four primary portals, namely, superolateral, superomedial, inferolateral and inferomedial. If necessary, additional portals were made to gain access to the posterior compartment. The basic instrument used consisted of a 5.5 mm full radius synovial resector, which was used in oscillating mode. Pressure irrigation was used for distension of the joint throughout the procedure using a gravity flow system.

Post-operatively, a compression bandage was applied leaving one or two portals open for drainage. The compression dressings were removed on the seventh post-operative day and progressive weight bearing started. The patients were placed on an exercise regimen to strengthen the quadriceps and hamstring muscles and to regain the range of motion. Standardised medical treatment was

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maintained throughout.

Patients were reviewed at 1, 3 and 6 months and after that every year. At each visit patients were scored for pain, range of motion, functional activity, and recurrent synovitis with effusion. The scores obtained under each criterion were evaluated using the modified score developed for evaluation of knee synovectomy, originally proposed in 1974 by Laurin [1].

Results and analysis

There were 46 patients, and six patients had bilateral involvement, making 52 knees, 26 of them were males. The ages ranged from 11 to 63 years (mean age 30.3). The minimum duration of pain was for 6 months and the maximum 60 (mean age 23.5). All 46 patients had pain and swelling, and 34 (74%) had reduced range of movement. The average hospital stay after operation was 1.8 days with a range of 1–8 days (Table 1).

Pain

During follow up, until the second year, an improvement in pain (from 88% to 96%) was noted, which was not statistically significant (p = 0.14). There was a significant worsening of pain in the following 3 years (from 96% to 78%), with p value of 0.01. But over a total 5 years of follow up, no significant worsening of pain (from 88.5% to 78%; p = 0.19) occurred.

Range of motion (ROM)

No significant loss in ROM in the first 2 years was noted. Over a period of 3 years there was a significant loss in ROM (p = 0.01), which worsened in the following years.

Functional activity

There was a significant loss of function 1 month after surgery (p = 0.001), which was completely regained in the next 2 months. At the end of 3 years there was a significant loss of function (from 100% to 90%) (p = 0.02) which correlates with worsening of pain and loss of ROM.

Synovitis and effusion

Eighty per cent of the patients showed improvement in achieving good or excellent results at the end of 1 month. The improvement increased to 100% at the end of 3 and 6 months, and then reduced to 96% at the end of 12 months and 2 years. The relatively small improvement at the end of 1 month was due to recurrent effusion post-operatively, which gradually settled at the end of 3 months. Recurrent synovitis was frequently noted during further follow up.

Discussion

Arthroscopic synovectomy has been developed as an effective tool for the ablation of the synovium in the knee joint. The procedure must be done in good time (early synovectomy) and be both radical and tissue sparing [2]. Effective and safe methods of access to different compartments of the knee have been developed. Providing that proper precautions be taken, the procedure can be carried out safely and with minimum morbidity [3]. In our study, 96% of the operated knees showed good results at the end of 6 months, that was maintained at the end of 12 months. This result gradually reduced to 90% and to 80% at the end of 4 and 5 years of follow up. This data compares favourably with the study of 25 cases by Smiley and Wasilewski (1990) [4]. They had 90% good results at the end of 2 years and 57% of the knees continued to show improvement after 4 years. Our study has shown that arthroscopic synovectomy is useful in relieving pain, and improving motion and functional activity. This procedure removes inflamed synovium and favours the local action of drugs (biomechanical indication) [5]. Our patients had a brief hospital stay of 1.8 days on an average. There was no need for post-operative manipulation under anaesthesia in any of the patients, and no functionally significant deterioration in range of motion at the end of 3 years. The maintenance of the benefits achieved and its later deterioration are on par with all previous studies [4].

A major success of arthroscopic synovectomy is its negligible rate of complications compared to open synovectomy [5]. Our series recorded no operative or post-operative complications with regard to bleeding, wound complications, neurovascular damage, post-operative pain and fibroarthrosis. Painless effusion was seen in 20% of patients up to 3 months post-operatively, that gradually settled. Radiation synovectomy after an interval of 6–8 weeks can be a good adjunct in the future to control recurrence and to treat the residual problems following arthroscopic synovectomy [3]. The repeatability of this technique because of its mild surgical insult gives it much advantage in treating recurrent cases.

Table 1. Number of knee joints with clinical criterion at each follow up (N = 52)

<table>
<thead>
<tr>
<th>Months of follow up</th>
<th>1</th>
<th>3</th>
<th>6</th>
<th>12</th>
<th>24</th>
<th>36</th>
<th>48</th>
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<tbody>
<tr>
<td>Improvement in pain</td>
<td>46</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>47</td>
<td>44</td>
<td>41</td>
</tr>
<tr>
<td>Improvement in ROM</td>
<td>52</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>47</td>
<td>45</td>
<td>42</td>
<td>39</td>
</tr>
<tr>
<td>Improvement in function</td>
<td>40</td>
<td>52</td>
<td>52</td>
<td>50</td>
<td>50</td>
<td>47</td>
<td>43</td>
<td>37</td>
</tr>
<tr>
<td>No recurrence of synovitis</td>
<td>42</td>
<td>52</td>
<td>52</td>
<td>50</td>
<td>50</td>
<td>42</td>
<td>37</td>
<td>28</td>
</tr>
</tbody>
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ROM = range of motion
Anti-tuberculosis drug induced hepatitis – a Sri Lankan experience

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(Index words: Liver function tests, WHO treatment recommendations)

Abstract

Objective To assess the incidence of anti-tuberculosis (TB) drug induced hepatitis (AIH) in Sri Lankan patients, determine risk factors of AIH, and to address management options in AIH.

Design A prospective study.

Setting Chest Hospital, Welisara, Sri Lanka, from April 2001 to April 2002.

Patients Seven hundred and eighty three patients with a confirmed diagnosis of TB and resident in the Colombo and Gampaha districts who presented to Chest Hospital, Welisara, Sri Lanka.

Methods WHO recommended treatment was commenced in all cases. AIH was diagnosed when patients complained of decreased appetite with nausea or vomiting and elevated serum bilirubin (SB; >1.1 mg/dL) or elevated serum alanine transferase (ALT; > 3 times upper limit of normal).

Results Of 783 enrolled patients, 74 (9.5%) developed AIH, the majority (58%) developing AIH within the first 2 weeks of the intensive phase of treatment. AIH was more common among patients over 60 years (p = 0.018), who developed pulmonary TB (p = 0.028), and in patients weighing 33–55 kg (p = 0.004). Age, weight and rifampicin overdosage were significant predictors of AIH. Of the 74 AIH patients, standard treatment was restarted in 60, treatment modified in six, two defaulted and six died.

Conclusions The incidence of AIH in Sri Lanka is 9.5% in treated patients. AIH was associated with age, low body weight and rifampicin overdosage.

References


Introduction

Anti-tuberculosis (TB) drug induced hepatitis (AIH) is a common complication in the management of TB. Studies in the USA and UK have reported a 3% and 4% incidence of AIH with rifampicin and isoniazid (with or without pyrazinamide in UK) [1,2], and studies from India have reported incidences ranging from 2% to as high as 30% [3–5]. No data are available for Sri Lanka.

Many risk factors for AIH have been described. They include advanced age, high alcohol consumption, extensive disease, hypoalbuminaemia, slow acetylator phenotype, female sex and endemic viral hepatitis [6–8]. The objective of this study was to assess the incidence of AIH in Sri Lankan patients, determine the risk factors of AIH and address management options.

Materials and methods

A prospective study was carried out from April 2001 to April 2002 at Chest Hospital, Welisara, Sri Lanka. Patients with a confirmed diagnosis of TB, resident in the Colombo and Gampaha districts and were under the care of the principal author were recruited after obtaining informed written consent. Diagnosis of TB was made according to WHO case definitions [9,10].

WHO category 1 and 2 treatment was commenced on all new and re-treatment cases respectively [9,10]. Baseline pretreatment serum bilirubin (SB) concentrations by the diazo method (normal range 0.2–1.1 mg/dL) and alanine transaminase (ALT) concentrations by the