Complete sectioning of axillary sentinel nodes in patients with breast cancer. Analysis of two different step sectioning and immunohistochemistry protocols in 246 patients

G Cserni

Aims: To evaluate two detailed step sectioning protocols for sentinel lymph nodes (SLNs).

Methods: After vital dye or combined dye and radiocolloid guided biopsy, SLNs were fixed in formalin and embedded in paraffin wax. In protocol A, SLNs from 123 patients were sectioned in steps of 50–100 µm, whereas in protocol B, SLNs from 123 patients were sectioned at steps of 250 µm. Epithelial marker immunohistochemistry (IHC) was performed on multiple levels in cases with negative haematoxylin and eosin findings.

Results: In groups A and B, 74 and 47 patients were found to have tumour cells in their axillary SLNs, and 19 (28%) and 18 (19%) patients, respectively, were upstaged as compared with the standard histological assessment. Nodal involvement detected by deeper sections was often micrometastatic or in isolated tumour cells.

Conclusions: Serial sectioning and IHC are recommended for the evaluation of SLNs. The optimal extent of the histopathological work up should be studied further.

The sentinel lymph node (SLN) theory in breast cancer suggests that the metastatic spread of mammary carcinomas through the lymphatic vessels follows an orderly pathway, and that only one or, at most, a few lymph nodes, the SLNs, are reached first by the process of metastasis. These are therefore the most likely sites of lymphogenic metastases. Identification, biopsy, and histological assessment of these nodes seems an ideal procedure for the qualitative staging of breast cancer, to clarify whether the tumour has spread to the axilla or not.1 It is generally believed that patients who have a positive SLN require axillary dissection to ensure a more quantitative staging, by indicating the number of lymph nodes involved. Axillary dissection can also provide a form of regional disease control, but if this option is not taken then treatment should include radiotherapy to the axilla. Alternatively, those patients who have negative SLNs are not likely to benefit from dissection or further treatment of the axilla.

“The sentinel lymph node theory in breast cancer suggests that the metastatic spread of mammary carcinoma through the lymphatic vessels follows an orderly pathway”

Because the SLNs are the most likely sites of regional metastasis, their histopathological assessment provides an opportunity for a more accurate staging by way of a more intensive investigation.1,2 However, there are many controversial issues regarding the extent of the histopathological work up, which lacks uniformity.3 Our study reports on the yields of two extensive protocols for the histopathological evaluation of axillary SLNs.

MATERIALS AND METHODS

The method of SLN biopsy (SLNB) first involved the peritumoral injection of patent blue dye, but a combined dye and radiocolloid guided technique preceded by lymphoscintigraphy is now preferred. Both tracers were administered peritumorally, although a few patients with negative lymphoscintigrams also received a smaller dose of radioactive tracer subareolarly. Technical details of the two biopsy methods, and their comparison, have been reported elsewhere.4,5 Briefly, blue lymphatics were always looked for first, and this was followed by a search for the remaining radioactive nodes. SLNs were defined as blue nodes, nodes at the end of a blue lymphatic, or radioactive nodes with an ex vivo radioactivity level > 10 times the background value. SLNs were sent for pathological assessment separately.

At the beginning of our study, which comprised our institutional validation phase, all patients (n = 191) but five underwent level I and II or complete axillary dissection after the SLNB. Four of these five patients had in situ carcinomas diagnosed preoperatively as malignant, but not as in situ carcinomas, and one had a cribriform carcinoma measuring 1 mm. Later, axillary dissection was planned only if the SLNs were positive (n = 55).

All SLNs were fixed in 7% buffered formalin. Smaller nodes were embedded in paraffin wax in toto, whereas larger nodes were bisected and both halves were processed.

In group A (protocol A), all of the SLNs were sectioned serially until the wax blocks were completely used up. Not all 3 to 5 µm thick sections were taken for histology, but every 10th to 20th section was examined (depending on nodal size) and, for a given node, the depths between the examined sections were approximately the same. This method of sectioning resulted in the examination of sections relatively equidistant (50–100 µm) from each other, depending on the overall nodal size, the distances being greater for larger SLNs. After every sixth section taken for haematoxylin and eosin (HE) staining, one was taken for cytokeratin immunohistochemistry (IHC).

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Abbreviations: HE, haematoxylin and eosin; IHC, immunohistochemistry; SLN, sentinel lymph node; SLNB, sentinel lymph node biopsy; VP, virtual protocol
(NCL-PAN-CK; Novocastra, Newcastle, UK; 1/100 dilution; or MNF116 (M0821); Dako, Glostrup, Denmark; 1/100 dilution). At the beginning of our study, epithelial membrane antigen (EMA; MU-182-UC; Biogenex, San Ramon, California, USA; 1/50 dilution) immunostaining was also performed in each node assessed, alternating with sections investigated by means of anticytokeratin antibodies. Immunostains were reported.

As reported previously, during the validation process of our study, epithelial membrane antigen (EMA; MU-182-UC; Biogenex, San Ramon, California, USA; 1/50 dilution) immunostaining was also performed in each node assessed, alternating with sections investigated by means of anticytokeratin antibodies. Immunostains were investigated only in the event of negative HE findings.

In group B (protocol B), the SLNs were sectioned at steps of 10–12 mm. These sections were stained with HE. Sections for IHC were taken after every third section stained, and were further processed in cases of negative HE findings.

A few obviously metastatic nodes were examined on only one or a few sections in both groups.

Tumours and metastases were classified on the basis of the TNM system. In the comparison of the two equally sized groups, the χ² test was used for categorical data, and the Student’s t test for continuous variables. Significance was set at p < 0.05.

**RESULTS**

Although the aim of our study was the description and possible comparison of two detailed histopathological analyses of SLNs, data on the results of SLNB itself are given here as background data required for a better understanding.

As reported previously, during the validation process of our study, the accuracy of the use of the SLNs to predict the overall axillary nodal status was 95.7% (176 patients with SLN positive). As part of this process, performance of the combined SLNB method on 72 patients during the validation phase yielded accuracy and false negative rates of 98.6% and 3.3%, respectively (one false negative SLNB/30 node positive patients).

In a previous report on the first 58 patients investigated by protocol A, a mean of 49 levels/SLN (range, 25–102) was reported. For protocol B, the mean number of levels/SLN was 21 (range, 7–46).

Table 1 compares the tumours of the patients in groups A and B. The tumours were significantly larger in the former group (χ² = 14.25; p < 0.05), their mean (SD) sizes being 2.30 (0.98) cm and 1.91 (0.90) cm, respectively (t = 3.15; p < 0.01).

Table 2 compares the metastases identified by the two protocols. Even when all the categories of table 2 were considered, there was a significant difference between the numbers of metastases identified in the two groups (χ² = 15.91; p < 0.025). It was mainly of interest to compare the metastatic yields of the two protocols, and accordingly the nodal status was taken as a positive result; the numbers of patients with positive SLNs/106 patients with a positive nodal status. As part of this process, performance of the combined SLNB method on 72 patients during the validation phase yielded accuracy and false negative rates of 98.6% and 3.3%, respectively (one false negative SLNB/30 node positive patients).

Table 1 shows the number of metastatic positive SLNs/106 patients with a positive nodal status. As part of this process, performance of the combined SLNB method on 72 patients during the validation phase yielded accuracy and false negative rates of 98.6% and 3.3%, respectively (one false negative SLNB/30 node positive patients).

Table 3 shows the figures when any nodal involvement was taken as a positive result; the numbers of patients with positive SLNs in a given tumour size category and their percentage of the total number of tumours in the same category are shown.
Figures 1 and 2 show the stepwise yields of sectioning protocols A and B, respectively.

Table 4 shows the proportion of isolated tumour cells, micrometastases, and macrometastases identified by different levels of the step sectioning protocols. It is evident from this table that metastases detected by deeper levels are more likely to be micrometastases or isolated tumour cells.

After analysing the real protocols, virtual protocols were also created on the basis of protocol A. In these virtual protocols (VPs) only every third (VP3), fourth (VP4), fifth (VP5), or sixth (VP6) level of the SLNs from group A were hypothetically examined. Table 5 shows the results of their analysis.

DISCUSSION

The identification of SLNs makes these nodes suitable for a more intensive histological work up. Serial sectioning and IHC would be too expensive for all axillary nodes, but these special methods may be cost effective for the detection of occult metastases in a limited number of specific nodes, the SLNs.

However, because most of our knowledge of the prognostic relevance of nodal metastases is based on standard pathological assessments, metastases detected by special methods are of uncertain biological relevance, and some authors recommend only standard assessment of the SLNs in the routine setting.6–8 Despite the fact that the biological meaning of minimal nodal involvement (micrometastatic disease and isolated tumour cell involvement) is unclear, there is growing evidence that their presence may indicate a somewhat worse outcome, at least in some subgroups of patients.9–13 Currently, most studies reporting on SLNB use special techniques to detect minimal involvement.14 However, the extent of the histopathological work up varies from study to study.15

“Many laboratories still use a single level assessment for the determination of the nodal status of malignant tumours, despite the fact that the inadequacy of such methodology was pointed out several decades ago.”16

In our present study, the yields of two extensive histological protocols for SLNs were assessed without considering the issue of the biological importance of the metastatic deposits identified. Because of the stricter inclusion criteria, the tumours studied later in the course of our SLNB series were smaller, and this could impact on the rate of their metastatic involvement; therefore, the results of the two methods used must be compared with caution. The rates of isolated tumour cells and micrometastases in groups A and B were 23% and 32%, respectively. This suggests that the difference in tumour size contributed considerably to the difference in the rate of nodal involvement identified, which is also reflected by the virtual protocols, where the upstaging rate of VP6 (21%) was above that of protocol B (19%), despite the larger steps in sectioning. When the nodal involvement detected in each tumour size category was considered (table 3), it became clear that the higher yield of protocol A was also present in most categories.

### Table 4

Proportion of isolated tumour cells and/or micrometastases and macrometastases to the total number of metastases detected by each protocol stratified by levels

<table>
<thead>
<tr>
<th>Protocol</th>
<th>Levels</th>
<th>Metastases detected</th>
<th>ITC</th>
<th>Micrometastases</th>
<th>ITC or micrometastases</th>
<th>Macrometastases</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>1–5</td>
<td>62</td>
<td>3 (5%)</td>
<td>4 (6%)</td>
<td>7 (11%)</td>
<td>55 (89%)</td>
</tr>
<tr>
<td></td>
<td>6–10</td>
<td>4</td>
<td>2 (50%)</td>
<td>2 (50%)</td>
<td>4 (100%)</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>11–15</td>
<td>2</td>
<td>1 (50%)</td>
<td>1 (50%)</td>
<td>2 (100%)</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>16–20</td>
<td>4</td>
<td>2 (50%)</td>
<td>2 (50%)</td>
<td>4 (100%)</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>&gt;20</td>
<td>2</td>
<td>1 (50%)</td>
<td>0</td>
<td>1 (50%)</td>
<td>1 (50%)</td>
</tr>
<tr>
<td>B</td>
<td>1</td>
<td>29</td>
<td>1 (3%)</td>
<td>4 (14%)</td>
<td>5 (17%)</td>
<td>24 (83%)</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>4</td>
<td>0</td>
<td>1 (25%)</td>
<td>1 (25%)</td>
<td>3 (75%)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>3</td>
<td>0</td>
<td>1 (33%)</td>
<td>1 (33%)</td>
<td>2 (67%)</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>2</td>
<td>1 (50%)</td>
<td>0</td>
<td>1 (50%)</td>
<td>1 (50%)</td>
</tr>
<tr>
<td></td>
<td>5</td>
<td>3</td>
<td>1 (33%)</td>
<td>1 (33%)</td>
<td>2 (67%)</td>
<td>1 (33%)</td>
</tr>
<tr>
<td></td>
<td>&gt;5</td>
<td>6</td>
<td>2 (33%)</td>
<td>3 (50%)</td>
<td>5 (83%)</td>
<td>1 (17%)*</td>
</tr>
</tbody>
</table>

*Metastasis first seen at the 6th level.
ITC, isolated tumour cells.

### Table 5

Analysis of virtual protocols derived from protocol A

<table>
<thead>
<tr>
<th>A, VP3</th>
<th>VP4</th>
<th>VP5</th>
<th>VP6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients with involved SLNs detected</td>
<td>74</td>
<td>72</td>
<td>70</td>
</tr>
<tr>
<td>Number of ITC undetected</td>
<td>1</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Number of micrometastases undetected</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Number of macrometastases undetected</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Number (% of patients upstaged)</td>
<td>19 (28)</td>
<td>17 (25)</td>
<td>15 (22)</td>
</tr>
</tbody>
</table>

A, protocol A; ITC, isolated tumour cells; VP3–6, virtual protocols examining only every 3rd, 4th, 5th, and 6th level of the sentinel lymph nodes (SLNs) from group A, respectively.
particularly in the pT1c and pT2 tumours, which were the most common ones found in the series. Because of the different tumour sizes in the two groups, the yields of the two histopathological protocols studied are best viewed separately.

Many laboratories still use a single level assessment for the determination of the nodal status of malignant tumours, despite the fact that the inadequacy of such methodology was pointed out several decades ago. Some of the recent recommendations consider a single section from each lymph node to be sufficient, although there are statements on the macrosectioning of larger nodes for better fixation and examination. The National Health Service breast screening programme in the UK recommends taking up to four separate examination.

As a method of ultrastaging in some studies, transcription polymerase chain reaction has also been applied to the identification of small metastases in SLNs and the study of their effect on non-SLN metastases. The size of the SLN metastasis to axillary dissection versus no further surgery) are available.

Both of our protocols identified a substantial proportion of the metastases in the first or the first few sections (figs 1, 2), and this is a general phenomenon seen in multiple studies assessing the role of a more intensive histopathological work up summarised in table 6. A single level HE stain approach would have identified somewhat more than 60–70% of the metastatic nodes. Although metastases in the SLNs show a predilection for the site where the tumour draining lymphatic vessel reaches the lymph node, the identification of this site requires vital dye labelling of the SLN, in addition to a piece of the blue stained lymphatic vessel. There has also been a suggestion by Turner et al that the site most likely to contain metastatic deposits is often located opposite the hilum of the node. Often, neither the inflow point of the tumour draining lymphatic vessel, nor the plane opposite to the hilum are easy

### Table 6: Review of studies assessing the role of serial or step sectioning and immunohistochemistry in the assessment of sentinel lymph nodes

<table>
<thead>
<tr>
<th>1st author</th>
<th>Number of patients</th>
<th>Number (%) of patients positive by standard HE</th>
<th>Protocols compared*</th>
<th>Number (%) of patients upstaged</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Jannink et al</td>
<td>19</td>
<td>6 (31.6%)</td>
<td>1 HE vs SS and IHC at 0.5 mm</td>
<td>3 (23.1%)</td>
<td>2 patients upstaged by SS and 1 by IHC</td>
</tr>
<tr>
<td>Kelley et al</td>
<td>28</td>
<td>9 (32.1%)</td>
<td>1 HE vs 4 level HE and 2 level IHC</td>
<td>2 (10.5%)</td>
<td>Evaluates SS too; distance of levels not stated</td>
</tr>
<tr>
<td>Czerniecki</td>
<td>41</td>
<td>12 (29.3%)</td>
<td>1 HE vs 4 level IHC</td>
<td>3 (10.3%)</td>
<td></td>
</tr>
<tr>
<td>Turner et al</td>
<td>52</td>
<td>10 (19.2%)</td>
<td>2 HE at 0.04 mm vs 2 further levels at 0.16 mm from each other</td>
<td>2 (4.8%)</td>
<td></td>
</tr>
<tr>
<td>Turner et al</td>
<td>52</td>
<td>10 (19.2%)</td>
<td>2 HE at 0.04 mm vs 2 IHC at 0.04 mm</td>
<td>8 (19%)</td>
<td></td>
</tr>
<tr>
<td>Turner et al</td>
<td>52</td>
<td>10 (19.2%)</td>
<td>2 HE at 0.04 mm vs 8 further levels IHC at 0.04 mm from each other</td>
<td>9 (21.4%)</td>
<td>Evaluates SS too</td>
</tr>
<tr>
<td>Naguchi et al</td>
<td>62</td>
<td>24 (38.7%)</td>
<td>1 HE vs some level IHC</td>
<td>1 (2.6%)</td>
<td>Retrospectively, the metastasis identified by IHC could have been seen on HE</td>
</tr>
<tr>
<td>Viola et al</td>
<td>155</td>
<td>45 (29%)</td>
<td>1 HE vs 14 further levels at 0.05 mm from each other; frozen sections</td>
<td>25 (16.1%)</td>
<td>IHC did not increase the sensitivity of SLN assessment</td>
</tr>
<tr>
<td>Pendás et al</td>
<td>478</td>
<td>93 (19.5%)</td>
<td>1 HE vs some level IHC</td>
<td>41 (10.6%)</td>
<td></td>
</tr>
<tr>
<td>Kowalczyk et al</td>
<td>33</td>
<td>8 (24.4%)</td>
<td>2 HE vs some level IHC</td>
<td>4 (16%)</td>
<td>2 pN1a, 2 pN0(i+)</td>
</tr>
<tr>
<td>Liu et al</td>
<td>38</td>
<td>12 (31.6%)</td>
<td>1 HE vs 3 further HE sections and IHC</td>
<td>5 (19.2%)</td>
<td>2 patients upstaged by HE and 3 by IHC; distance of levels not stated</td>
</tr>
<tr>
<td>Nährig et al</td>
<td>40</td>
<td>18 (45%)</td>
<td>1 HE vs 4 further level HE at 0.15 mm from each other</td>
<td>4 (18.2%)</td>
<td></td>
</tr>
<tr>
<td>Mann et al</td>
<td>51</td>
<td>10 (19.6%)</td>
<td>1 HE vs some level IHC</td>
<td>10 (24.4%)</td>
<td></td>
</tr>
<tr>
<td>Weaver et al</td>
<td>386</td>
<td>104 (27.0%)</td>
<td>1 HE vs 2 further HE levels at 0.1 mm from each other</td>
<td>19 (8.9%)</td>
<td></td>
</tr>
<tr>
<td>Pelayo et al</td>
<td>68</td>
<td>21 (30.9%)</td>
<td>1 HE vs SS with IHC only at 0.25 mm</td>
<td>12 (25.5%)</td>
<td></td>
</tr>
<tr>
<td>Dowlatshahi et al</td>
<td>200</td>
<td>34 (17%)</td>
<td>1 HE vs SS with IHC only at 0.25 mm</td>
<td>51 (30.7%)</td>
<td>24 pN1a, 27 pN1o(i+)</td>
</tr>
<tr>
<td>Torregrosa et al</td>
<td>250</td>
<td>69 (27.6%)</td>
<td>1 HE vs 4 further HE at 0.25 mm from each other</td>
<td>17 (5.9%)</td>
<td></td>
</tr>
<tr>
<td>Torregrosa et al</td>
<td>250</td>
<td>69 (27.6%)</td>
<td>1 HE vs 4 further level HE at 0.25 mm from each other</td>
<td>17 (9.4%)</td>
<td>Evaluates SS too</td>
</tr>
<tr>
<td>Wong et al</td>
<td>973</td>
<td>104 (10.7%)</td>
<td>1 HE vs 2 level IHC</td>
<td>58 (6.7%)</td>
<td></td>
</tr>
<tr>
<td>Yarei et al</td>
<td>96</td>
<td>0</td>
<td>1 HE vs 2 level HE and 1 level IHC at 0.005 mm from each other</td>
<td>17 (17.7%)</td>
<td></td>
</tr>
<tr>
<td>This study A</td>
<td>123</td>
<td>55 (44.7%)</td>
<td>1 HE vs SS at 0.05–0.1 mm, IHC levels at 0.3–0.6 mm from each other</td>
<td>19 (27.9%)</td>
<td>1 pN1a and 3 pN0(i+) first identified by IHC</td>
</tr>
<tr>
<td>This study B</td>
<td>123</td>
<td>29 (23.6%)</td>
<td>1 HE vs SS at 0.25 mm, IHC levels at 0.75 mm from each other</td>
<td>18 (19.1%)</td>
<td>3 pN1a and 3 pN1o(i+) first identified by IHC</td>
</tr>
</tbody>
</table>

*The standard HE serving as baseline examination comprised halving or macrosectioning SLNs at 2–3 mm and examining 1 HE section from each part, except in 2 studies, where 2 HE sections were obtained.

HE, haematoxylin and eosin; IHC, immunohistochemistry against epithelial markers, generally cytokeratin with AE1/AE3, MIB-1, CAM 5.2, PanCK antibodies; some studies also used epithelial membrane antigen; pN1a, micrometastasis; pN0(i+), isolated tumour cells; SS, serial sectioning.
thought that a more accurate estimate of upstaging by IHC might be as high as 17% after examining five levels at the central cross section (group B in our study), the chances of detecting macrometastases (>2 mm), whereas subsequent levels are more likely to identify micrometastases or isolated tumour cells. As far as we are aware, no study has given details on the addition of serial sectioning, the role of IHC in identifying metastatic cells or micrometastases detected and the lower the value of small sectioning steps, the higher the rate of metastatic small metastatic foci in SLNs on non-SLN involvement and on the extent of step sectioning and IHC must be balanced against each other to attain a reasonable assessment protocol for the work up of sentinel lymph nodes. Although metastases are not uncom­monly distributed away from the central cross section, these studies are those that document the highest rate of conversion from standard HE negative SLNs to involved SLNs. With the use of extensive step sectioning, the role of IHC in identifying metastastic cells or groups of metastastic cells decreases; IHC identified only 5.4% and 12.8% of the involved nodes in our protocols A and B, respectively, whereas it helped to strengthen the suspicion of metastastic cells in only three cases in the Milan series. The smaller the sectioning steps, the higher the rate of metastastic cells or micrometastases detected and the lower the value of IHC.

The combination of step sectioning and IHC gives the best rates of identification of occult metastatic deposits. The costs and affordability of step sectioning and IHC must be balanced against each other to attain a reasonable assessment protocol for the work up of SLNs. It must be accepted that histopathology is based on tissue sampling and has a rate of unidentified lesions (including occult metastases). If it is assumed that the identification of an isolated tumour cell and a group of 10 tumour cells randomly placed in a 1 cm SLN would require 312 and 139 sections, respectively, each of the SLN histology protocols studied to date must have missed a few cases of minimal nodal involvement of unknown importance. Complete examination of any routinely assessed specimen is not feasible, but a compromise between workload, costs, and sensitivity should be found. Our study is one step towards finding such a compromise. At least, it can be concluded from our results that the combination of step sectioning and IHC may be recommended for the work up of SLNs. Whether the detection of small metastatic foci (isolated
tumour cells or micrometastases) detected in deeper sections is clinically relevant or not is not known. As discussed previously, the size of SLN metastases might be important in either predicting further nodal involvement and therefore indicating some type of regional treatment) or in influencing survival and prognosis (and therefore influencing the indication of adjuvant systemic treatment). SLNB has a recognised false negative rate, which seems acceptable to many. A certain risk of further nodal involvement may also be acceptable, in addition to a minor prognostic disadvantage. Therefore, the extent of step sectioning and the impact of extremely small metastatic foci in SLNs on non-SLN involvement and on the fate of the patients remain to be elucidated in prospective clinical trials and probably cost benefit analyses.

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REFERENCES
Enzymes aid invasion of arthritic joints

Recent evidence suggests that enzymes which destroy cartilage aid invasive growth of fibroblast-like synoviocytes (FLS) in rheumatoid arthritis (RA). An in vitro study has compared potential invasive properties of FLS from patients with RA, osteoarthritis (OA), and avascular necrosis (AVN) and found that FLS from RA had significantly more matrix metalloproteinases (MMPs).

Growth through an artificial matrix was greater for FLS from RA (median cell number 4788 v 1875 for OA, v 1350 for AVN), and so was growth rate (0.27/day v 0.22/day v 0.25/day, respectively). However, growth rate showed no correlation with cell number. FLS expressing MMP-1, MMP-3, or MMP-10 were significantly more invasive (median number of invasive cells 3835, 4248, and 4990, respectively) whether from RA or OA. But the odds of having MMP-1 and MMP-9 and RA were significant, 6.5 and 10.7, when compared with OA. Other attributes—expression of cathepsin-K and tissue inhibitors of MMP-1 and MMP-2—did not influence invasiveness. FLS were cultured from tissue obtained from joint replacements or synovectomy in patients with RA (30), OA (17), and AVN (nine). Invasiveness was assayed in a Matrigel transwell culture system, by counting cells that migrated through the matrix after three days’ incubation. Growth rate was determined from cell counts of cultures harvested at intervals after seeding. Expression of cathepsin-K, tissue inhibitors, and MMPs was indicated by reverse transcriptase-PCR.

Activated FLS invade the synovium, articular cartilage, and bone in RA. Whether this is through increased growth or invasiveness has not been studied directly before now.

### References


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**Enzymes aid invasion of arthritic joints**

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Complete sectioning of axillary sentinel nodes in patients with breast cancer. Analysis of two different step sectioning and immunohistochemistry protocols in 246 patients

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- Breast cancer (506)
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