Optimal excision margins for primary cutaneous melanoma: a systematic review and meta-analysis

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Objective: To determine in patients with localized primary melanoma of the trunk or extremities the optimal excision margin that achieves the highest disease-free survival and overall survival and the lowest local recurrence rate. Data sources: Trials comparing 2 different excision margins were identified by searching MEDLINE from 1966 to May 2002 using the term “melanoma,” subheading “surgery,” and limiting the search to human studies and randomized controlled trials (RCTs). Additional studies were found using the MeSH term “surgical procedures, operative,” combining with “melanoma,” and limiting to human studies. We searched EMBASE and the Cochrane Library in May 2002 using similar terminology. No language restriction was applied. Study selection: We selected studies for the overview using the following inclusion criteria: design — an RCT with wide excision versus narrower excision (margin width was not specified a priori); population — adult patients (> 18 yr) with cutaneous melanoma of the trunk or extremities without evidence of metastasis; intervention — surgical excision of the primary melanoma; and outcomes — at least 1 of overall survival, disease-free survival, local recurrence, wound complications and necessity for skin grafting. Data extraction: Information was abstracted for each outcome reported in the studies, and results were pooled by consensus. Statistical analysis was performed using RevMan 4.1 (The Cochrane Collaboration) software program. Relative risk and risk difference were reported with 95% confidence intervals. The number needed to harm was calculated for the need for skin grafting by taking the inverse of the risk difference. Data synthesis: Three trials and their follow-up studies met the inclusion criteria and included 2087 adults with localized cutaneous melanoma of the trunk or extremities. No statistically significant differences were found between wide surgical excision (margins ranging from 3–5 cm) and narrower surgical excision (margins ranging from 1–2 cm) with respect to mortality, disease-free survival or local recurrence rate. Conclusions: Surgical excision margins no more than 2 cm around a melanoma of the trunk or extremities are adequate; overall survival, disease-free survival and recurrence rate are not adversely affected compared with a wider excision. There is more data to support a 2-cm margin than a 1-cm margin as the minimum margin of excision. Surgical margins should be no less than 1 cm around the primary melanoma.

Surgical excision remains the mainstay of treatment for primary cutaneous melanoma. However, how much surrounding normal skin should be excised around a primary cutaneous melanoma is controversial. Historically, specimens larger than 5 cm in dimension were removed because of the contention that a large excision will provide optimal outcomes in terms of local recurrence rate and survival. Furthermore, because increased numbers of melanocytes have been found in the clinically normal skin surrounding many melanomas and melanocytic activation has been reported up to 4 cm from the primary tumour in clinically normal skin, wide excision is a logical choice. In some cases, extremely wide excisions were recommended because recurrences were found up to 15 cm from the original site of primary tumour excision. Acceptance of such a wide surgical margin was not based on good scientific outcome data, and these radical excisions were morbid, especially since many required skin grafting.

Other studies have suggested that more conservative excision margins are equivalent to large excisions with respect to survival yet allow a much less morbid procedure. In addition, Olsen reported similar local recurrence rates in patients who had a narrow excision and those who had at least 5 cm excision margins. The suggestion from these nonrandomized studies that narrower excisions are equivalent to wider excisions can only be confirmed in randomized controlled trials (RCTs). A major advantage of narrower excisions, if overall survival and recurrence rates are similar, would be less morbidity and fewer skin grafting procedures, which require the use of additional resources, more operative time, more intensive wound care, and usually have a less desirable cosmetic outcome.

Our primary objective was to determine the optimal margin of surgical excision that achieves the highest disease-free survival, overall survival and the lowest local recurrence rate in adult patients with a primary cutaneous melanoma of the trunk or extremities and no evidence of metastatic disease. Secondary objectives were to determine the complication rate in terms of local wound infection and dehiscence rate, and the need for skin grafting with various margins of excision.

Methods

This systematic review and meta-analysis was completed using the approach described by The Cochrane Collaboration.

Data sources

All RCTs or quasi-randomized clinical trials comparing 2 different margins of excision in adult patients with confined primary cutaneous melanoma of the trunk or extremities were searched on MEDLINE from 1966 to May 2002 using the terms “melanoma, surgery,” limited to human studies. Additional studies were found using the MeSH term “Surgical Procedures, Operative,” combined with “Melanoma,” and limited to human studies. EMBASE and the Cochrane Library were searched in May 2002 using similar terminology. No language restriction was applied. No unpublished articles or abstracts published or presented at meetings were included.

Study selection

Two authors (P.I.H. and L.A.D.) reviewed the titles of the articles and the abstracts if available as identified from the search. Any article deemed appropriate or possibly meeting the inclusion criteria from the title or abstract was retrieved. These articles were screened by both authors, and the following criteria were used to select studies: (1) study design — an RCT with one excision margin versus another excision margin (there were no strict a priori definitions of wide or narrow margins; any study that compared one excision margin width versus another automatically met the criterion of a wide versus a narrower excision margin); (2) population — adult patients (≥ 18 yr) with cutaneous melanoma of the trunk or extremities without evidence of metastasis; (3) intervention
— surgical excision of the primary melanoma; (4) outcome — at least 1 of overall survival, disease-free survival, local recurrence, wound complications and need for skin grafting. Studies were excluded if they were repeat reports of a previously published trial.

**Data extraction**

Data forms were developed and used to extract data from each article that met the inclusion criteria. The 2 authors independently abstracted the data for each article. Demographic data and the main outcome measures were extracted. Overall survival and disease-free survival were converted to mortality and any recurrence, in keeping with The Cochrane Collaboration format of attaining a risk reduction of an adverse event rather than of a favourable event.

Studies were assessed for methodologic quality using the “Users’ guides to the medical literature” published by the Journal of the American Medical Association.10,11

**Statistical analysis**

Analysis was performed using RevMan 4.1 (The Cochrane Collaboration, Oxford, UK) software program. Relative risk (RR) and risk difference (RD) are reported using the fixed-effects model with 95% confidence intervals (CIs). An RR or equal to 1 or an RD equal to 0 indicates no difference in outcome between wide excision and narrower excision. An RR less than 1 or a negative RD indicates that the outcome favours wider excision, and RR greater than 1 or a positive RD indicates that the outcome favours narrower excision. Differences in outcomes were considered statistically significant if the 95% CI does not include 1 for RR or 0 for RD. In the case of a statistically significant risk reduction or increase, the number needed to treat (or harm) was calculated. Two separate meta-analyses of mortality, any recurrence and local recurrence were performed using the original trials in one analysis and the follow-up studies in the second analysis. Mortality, any recurrence and local recurrence were grouped over a range of follow-up time because of different follow-up in each trial.

**Data synthesis**

**Description of studies**

Details of the included studies are shown in Table 1.12–17 Three studies totalling 1867 patients met the inclusion criteria. Two studies were international multicentre studies with participating centres in Europe, North and South America, and Russia.12,13 The third study was a multicentre study completed in Sweden.14 Three studies, all follow-up studies to the original 3, also met the criteria; the study by Cohn-Cedermark and associates17 added patients to the

**Table 1**

| Details of the 3 Randomized Clinical Trials (RCTs) (Primary and Follow-Up Studies) of Patients With Malignant Melanoma of the Extremities Who Underwent Wide or Narrower Excision |
|-------------|-----------------|-----------------|-----------------|-------------------|
| Series | Methods | Participants | Interventions | Outcomes |
| Veronesi et al, 198812 (primary study) | Multicentre international RCT | Blinding of: randomization: yes intervention: no outcome measures: no Complete follow-up: no (1 lost) | 612 adults with cutaneous melanoma up to 2 mm Breslow thickness | 307 patients treated with 3-cm margin v. 305 with 1-cm margin of excision. Age, sex, Breslow thickness, ulceration, anatomic site similar | 55 mo OS, DFS and LRR |
| Veronesi et al, 199113 (follow-up study) | Follow-up to Veronesi et al 1988 primary study | Same as Veronesi et al 1988 | Same as Veronesi et al 1988 | 8 yr OS, DFS and LRR |
| Balch et al, 199315 (primary study) | Multicentre, international RCT | Blinding of: randomization: not stated intervention: no outcome measures: only for local recurrence Complete follow-up: no (2.3% lost) | 486 adults with cutaneous melanoma 1–4 mm Breslow thickness | 242 patients treated with 4-cm margin v. 244 with 2-cm margin of excision. Age, sex, Breslow thickness, ulceration, anatomic site similar | 5 yr OS, median 6 yr LRR Wound complications Need for skin graft LOHS |
| Balch et al, 200116 (follow-up study) | Follow-up to Balch et al 1993 primary study | Same as Balch 1993 | Same as Balch et al 1993 | 10 yr mean OS and LRR |
| Ringborg et al, 199014 (primary study) | Multicentre, international RCT | Blinding of: randomization: yes intervention: no outcome measurements: can’t tell for recurrence Complete follow-up: yes | 769 adults with cutaneous melanoma 0.8–2 mm Breslow thickness | 396 patients treated with 5-cm margin v. 373 with 2-cm margin of excision. Age, sex, Breslow thickness, ulceration, anatomic site similar | 5 yr OS, 5 yr melanoma survival median 4 yr DFS, median 4 yr LRR |
| Cohn-Cedermark et al, 200017 (follow-up study) | Follow-up to Ringborg et al 1996 primary study with added patients | 989 adults with cutaneous melanoma | 513 patients treated with 5-cm margin v. 476 with 2-cm margin of excision | 11 yr OS and melanoma survival 8 yr DFS and LRR |

LOHS = length of hospital stay, OS = overall survival, LRR = local recurrence rate, DFS = disease-free survival.
original trial, which increased the total number to 2087. Three studies were excluded because they were repeat reports of the original studies.18–20

The trial of Veronesi and colleagues12 enrolled patients with melanomas up to 2 mm in Breslow thickness. Patients were randomized to wide excision of 3 cm or to 1-cm excision. After the skin incision was made, the flaps were undermined radially for at least 1 cm and continued to fascia. The 3 outcome measures were overall survival, disease-free survival and local recurrences. Local recurrence was defined as a first recurrence within the scar or within 1 cm from the scar.15 Mean duration of follow-up was 55 months. In the follow-up study had a mean duration of follow-up of 8 years.15

In the trial reported by Balch and associates,13 patients were enrolled if the melanoma was between 1 and 4 mm thick according to Breslow's classification. Patients were randomized to wide excision of 4 cm or 2 cm. Excisions incorporated the skin at the stipulated margin and underlying subcutaneous tissues to fascia; inclusion of fascia in the specimen was optional. This trial also had a second randomization to an elective lymph-node dissection or observation of the regional nodal basin. The outcome of local recurrence was defined as a biopsy-proven first recurrence within 2 cm of the scar; if the local recurrence occurred after the appearance of distant metastases, for example, then a local recurrence was not counted. Overall survival was another outcome measure. Median survival was 72 months. In the follow-up study the mean survival was 10 years.16

In their trial, Ringborg and colleagues14 enrolled 769 patients with melanoma ranging from 0.8 to 2.0 mm in Breslow thickness. Patients were randomized to wide excision of 5 cm or 2 cm. The excision technique was similar to that of Balch and associates.13 Outcome measures included overall survival, disease-free survival and local recurrence rate. Local recurrence was defined as recurrence within the scar or skin graft. Median follow-up was 5.8 years for survival, and 4 years for recurrences. The follow-up study included additional patients to a total of 989, and median follow-up of 11 years for survival and 8 years for recurrences.17

Study quality

In general, the methodologic quality of the 3 trials was good, recognizing that in surgical trials it is more difficult to blind the treating clinicians and physicians responsible for measuring the outcome. Therefore, because all studies were surgical, each lacked at least 1 important methodologic criterion that would possibly produce bias: they all lacked blinding of the intervention. In all trials it was difficult to determine whether individual validity criteria were met because often these aspects were not stated explicitly in the methods section. Particularly with respect to outcome measurements other than the dichotomous outcome of “alive” or “dead,” in these surgical trials the outcome of disease-free survival or local recurrence was often assessed by clinicians who may not have been blinded to treatment allocation. It is possible, therefore, that the clinicians responsible for detecting recurrences could introduce bias by measuring the outcome in a different manner, depending on their preference, if they knew the patient’s treatment allocation.

The trial sponsored by Veronesi and colleagues12 provided explicit details on randomization: the process was concealed. Only 1 patient was lost to follow-up from the wide excision arm. Analysis was not using the intention-to-treat principle, because those who received the wrong treatment (8 patients from the narrow excision group and 7 from the wide excision group) were excluded. The study of Balch and associates13 did not report whether allocation was concealed. We can assume that the intervention was known by participating surgeons and other clinicians. The outcome that was identified by a panel blinded to treatment was local recurrence: the panel of pathologists identified biopsy-proven recurrences, which would help to avoid bias for this outcome. Not stated is whether disease-free survival was assessed in a blinded fashion. Finally, although follow-up was not complete, only 2.3% were lost to follow-up. The actual number of patients lost in each arm was not stated.

In the Ringborg trial14 the methodology of randomization was outlined more clearly in the updated trial.17 A telephone system was used for randomization at central cancer centres, and presumably allocation would be blinded to enrolling physicians. Regarding outcomes of local recurrence and any recurrence, the authors did not state about blinding using adjudication panels, so it can be assumed that treatment allocation might have been deduced by the outcome assessors. Follow-up was complete.

Mortality

All 3 initial studies reported on mortality at 4 to 6 years’ follow-up (Fig. 1). Balch and associates13 showed an RR of death of 0.79 in favour of wide excision, but the 95% CIs ranged from 0.54 to 1.15, not statistically significant. The Veronesi12 and Ringborg14 trials showed a slight advantage of a narrower excision, with RRs of 1.02 and 1.19, respectively, but with 95% CIs ranging from 0.71 to 1.45 and 0.52 to 2.72, respectively. With all studies combined, the RR of death with wide excision compared to a narrower excision was not significantly decreased (RR = 0.93, 95% CI 0.73–1.19; RD = –0.01; 95% CI −0.04 to 0.02) (Fig. 1).

Mortality at 8 to 11 years’ follow-up was not significantly different between wide excision and a narrower ex-
cision (RR = 0.95, 95% CI 0.81–1.12; RD = −0.01; 95% CI −0.05 to 0.02) (Fig. 2).

**Any recurrence**

The trials of Veronesi and colleagues\(^\text{12}\) and Ringborg and colleagues\(^\text{14}\) reported any recurrence that occurred between 55 and 69.6 months (Fig. 3). Neither trial showed a significant difference between wide excision and a narrower excision. Also, the 2 studies together show no difference between wide versus narrower excision (RR 1.03, 95% CI 0.81–1.32; RD = 0.00, 95% CI −0.03 to 0.04) for any recurrence (Fig. 3).

At longer follow-up of 8 years, no difference (RR = 0.89, 95% CI 0.72, 1.09; RD = −0.02; 95% CI −0.06, 0.02) remained between the 2 interventions from the follow-up studies (Fig. 4).

**Local recurrence rates**

All 3 initial studies reported local recurrence rates; none showed differences in local recurrence rates between wide and narrower excisions at 48 to 72 months’ follow-up (Fig. 5). The Veronesi\(^\text{12}\) and Balch\(^\text{13}\) trials reported local recurrences only as the first recurrence, whereas the Ringborg trial\(^\text{14}\) reported total local recurrences. Combining the 3 studies revealed no difference in local recurrence between wide and narrower excisions (RR = 0.98, 95% CI 0.38–2.52; RD = 0.00, 95% CI −0.01, 0.01) (Fig. 5).

At 8 to 10 years, the local recurrence rates remained similar between interventions in the 3 follow-up studies (Fig. 6). Relative risk for local recurrence with the 3 follow-up studies combined was 0.90 (95% CI 0.41–2.00), and the RD was 0 (95% CI −0.01 to 0.01).

**Wound complications**

Only Balch and associates\(^\text{13}\) reported wound complications. Postoperative

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**Excision margins for primary cutaneous melanoma**

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**FIG. 1.** Comparison of mortality at 4 to 6 years between wide and narrower excision in the 3 original trials of melanoma of the extremities.

<table>
<thead>
<tr>
<th>Series</th>
<th>Wide excision no./total no.</th>
<th>Narrower excision no./total no.</th>
<th>Risk ratio (95% CI fixed)</th>
<th>Weight, %</th>
<th>Risk ratio (95% CI fixed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Balch et al, 1993(^\text{13})</td>
<td>39/242</td>
<td>50/244</td>
<td></td>
<td>44.7</td>
<td>0.79 (0.54–1.15)</td>
</tr>
<tr>
<td>Ringborg et al, 1996(^\text{14})</td>
<td>54/396</td>
<td>50/373</td>
<td></td>
<td>46.3</td>
<td>1.02 (0.71–1.45)</td>
</tr>
<tr>
<td>Veronesi et al, 1988(^\text{12})</td>
<td>12/307</td>
<td>10/305</td>
<td></td>
<td>9.0</td>
<td>1.19 (0.52–2.72)</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>105/945</td>
<td>110/922</td>
<td></td>
<td>100.0</td>
<td>0.93 (0.73–1.19)</td>
</tr>
</tbody>
</table>

Test for heterogeneity $\chi^2 = 1.34$, df = 2, $p = 0.51$. Test for overall effect $z = −0.58$, $p = 0.6$.

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**FIG. 2.** Comparison of mortality at 8 to 11 years between wide and narrower excision in the 3 follow-up trials of melanoma of the extremities.

<table>
<thead>
<tr>
<th>Series</th>
<th>Wide excision no./total no.</th>
<th>Narrower excision no./total no.</th>
<th>Risk ratio (95% CI fixed)</th>
<th>Weight, %</th>
<th>Risk ratio (95% CI fixed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Balch et al, 2001(^\text{16})</td>
<td>56/242</td>
<td>730/244</td>
<td></td>
<td>32.1</td>
<td>0.77 (0.57–1.04)</td>
</tr>
<tr>
<td>Cohn-Cedermark et al, 2000(^\text{17})</td>
<td>134/513</td>
<td>117/476</td>
<td></td>
<td>53.7</td>
<td>1.06 (0.86–1.32)</td>
</tr>
<tr>
<td>Veronesi et al, 1991(^\text{15})</td>
<td>30/307</td>
<td>32/305</td>
<td></td>
<td>14.2</td>
<td>0.93 (0.58–1.49)</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>220/1062</td>
<td>222/1025</td>
<td></td>
<td>100.0</td>
<td>0.95 (0.81–1.12)</td>
</tr>
</tbody>
</table>

Test for heterogeneity $\chi^2 = 2.87$, df = 2, $p = 0.24$. Test for overall effect $z = −0.60$, $p = 0.5$.
wound infection favoured wide excision (RR = 0.85, 95% CI 0.39–1.87; RD = –0.01, 95% CI –0.05 to 0.03) as did wound dehiscence (RR = 0.92, 95% CI 0.40–2.12; RD = 0, 95% CI –0.04 to 0.03), but these RRs and RDs were not statistically significant.

**Need for skin grafting**

Once again, only Balch and associates\(^1\) reported on the need for skin grafting. There was a significant reduction in the need for skin grafting with narrow excision compared with wide excision (RR = 4.15 95% CI 2.83–6.07; RD = 0.35, 95% CI 0.27–0.42; \(p < 0.001\)). The number needed to harm from wide excision is 3, with 95% CI ranging from 2.38 to 3.7; for every 3 patients who undergo wide excision, 1 patient will require a skin graft who would not have needed it if a narrower excision had been performed.

**Discussion**

This meta-analysis showed no benefit for a wide excision over a narrower excision in patients with melanoma of the trunk or extremities. The data indicate that an excision margin of no greater than 2 cm is adequate for treatment of primary melanoma, and that overall survival, disease-free survival and local recurrence rates are not adversely affected by this margin of excision.

The trials differed in eligibility criteria, particularly the criterion of Breslow thickness. The margins that were compared and the outcome measures differed. For these reasons, many questions remain unanswered, particularly the optimal narrower margin and whether different subsets of Breslow thickness can be treated with different excision margins. For instance, the surgical excision performed by Veronesi and colleagues\(^1\) differed from that of the other 2 studies: flaps were raised in all directions, and at the fascial level, the subcutaneous tissue in the narrow 1-cm excision specimen would have been similar to the 2-cm excision in the other trials. Local recurrences were defined differently in terms of distance from the scar and in terms of initial recurrence. Nevertheless, the events were very rare, and the excision margin had little impact on the

**FIG. 3.** Comparison of any recurrence at 55 to 70 months between wide and narrower excision in 2 original trials of malignant melanoma of the hands and feet. The trial reported by Balch and associates\(^1\) did not report this outcome.

<table>
<thead>
<tr>
<th>Series</th>
<th>Wide excision no./total no.</th>
<th>Narrower excision no./total no.</th>
<th>Risk ratio (95% CI fixed)</th>
<th>Weight, %</th>
<th>Risk ratio (95% CI fixed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ringborg et al, 1996(^1)</td>
<td>81/396</td>
<td>76/373</td>
<td></td>
<td>75.0</td>
<td>1.00 (0.76–1.33)</td>
</tr>
<tr>
<td>Veronesi et al, 1988(^1)</td>
<td>29/307</td>
<td>26/305</td>
<td></td>
<td>25.0</td>
<td>1.11 (0.67–1.84)</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>110/703</td>
<td>102/678</td>
<td></td>
<td>100.0</td>
<td>1.03 (0.81–1.32)</td>
</tr>
</tbody>
</table>

Test for heterogeneity \(\chi^2 = 0.11, df = 1, p = 0.74\).

Test for overall effect \(z = 0.24, p = 0.8\).

**FIG. 4.** Comparison of any recurrence at 8 years between wide and narrower excision in 2 follow-up trials of malignant melanoma of the extremities. The trial reported by Balch and associates\(^1\) did not report this outcome.

<table>
<thead>
<tr>
<th>Series</th>
<th>Wide excision no./total no.</th>
<th>Narrower excision no./total no.</th>
<th>Risk ratio (95% CI fixed)</th>
<th>Weight, %</th>
<th>Risk ratio (95% CI fixed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cohn-Cedermark et al, 2000(^1)</td>
<td>96/513</td>
<td>98/476</td>
<td></td>
<td>64.4</td>
<td>0.91 (0.71–1.17)</td>
</tr>
<tr>
<td>Veronesi et al, 1991(^1)</td>
<td>48/307</td>
<td>56/305</td>
<td></td>
<td>35.6</td>
<td>0.85 (0.60–1.21)</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>144/820</td>
<td>154/781</td>
<td></td>
<td>100.0</td>
<td>0.89 (0.72–1.09)</td>
</tr>
</tbody>
</table>

Test for heterogeneity \(\chi^2 = 0.09, df = 1, p = 0.77\).

Test for overall effect \(z = -1.13, p = 0.3\).
propensity for local recurrence.

From the results of this meta-analysis, it is difficult to recommend the minimum excision margin required. The minimum margin studied was 1 cm; however, more data were available on a 2-cm margin with 2 studies combined. The 2-cm excision margin has never been compared directly to a 1-cm margin in a randomized trial.

It has been proposed that a 1-cm margin of excision be performed for melanomas less than 1 mm thick, reserving wider excisions for tumours 1 mm or thicker.21,22 There are many descriptive studies supporting this policy, showing that the local recurrence rates are acceptable and similar to that of wider excisions. However, it should be noted that this meta-analysis does not strictly support this policy because the 3 major RCTs contain data on fewer than 185 patients with melanomas less than 1 mm thick who had a 1-cm excision. Therefore, a 2-cm margin should be the goal.

Only the Balch trial 13 reported on wound complications and the need for skin grafting. Total wound complications did not differ significantly between the wide and narrow excision. The need for skin grafting, not surprisingly, was higher in the wide excision group. However, only about 10% of patients required skin grafting with a 4-cm excision. Unfortunately, skin grafting requires more intense nursing and wound care than an excision that is closed primarily. With a number needed to harm of 3 with wide excision, a narrower excision readily helps to avoid the morbidity of a skin graft.

Conclusions

A surgical excision margin of no more than 2 cm around a melanoma of the trunk or extremities is adequate; mortality and recurrence rates should not be adversely affected compared to a wider margin. Surgi-
cal margins should be no less than 1 cm around the primary melanoma. A 1-cm margin has never been tested against a 2-cm margin in any randomized trial, but more data are available for a 2-cm excision as the minimum surgical margin for optimal overall survival, disease-free survival and local recurrence.

A trial that randomized patients with cutaneous melanoma up to 4 mm thickness to a 1-cm versus 2-cm surgical excision would answer the question of what is the optimal narrow excision. However, with such low event rates, and the fact that 90% of wounds probably could be closed primarily even with a 2-cm margin, the answer may be more academic than clinically useful. With such little impact of surgical margin on prognosis, in contrast to the great influence of tumour thickness, more effort should be expended on primary and secondary prevention so that early detection will decrease the proportion of patients presenting with thicker melanomas.

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Competing interests: None declared.

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