Table SI. Summary of the cases of cutaneous B-cell lymphoprolifera-tive disorders associated with anetoderma (1995–2013, ref (2–6))

<table>
<thead>
<tr>
<th>Number of patients</th>
<th>16</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex: male/female</td>
<td>14/2</td>
</tr>
<tr>
<td>Mean age (range)</td>
<td>44.8 (26–70)</td>
</tr>
<tr>
<td>Primary diagnosis</td>
<td></td>
</tr>
<tr>
<td>Cutaneous lymphoid hyperplasia</td>
<td>1</td>
</tr>
<tr>
<td>Cutaneous B-cell lymphoma</td>
<td>4</td>
</tr>
<tr>
<td>Cutaneous plasmacytoma</td>
<td>1</td>
</tr>
<tr>
<td>Primary cutaneous marginal zone lymphoma</td>
<td>7</td>
</tr>
<tr>
<td>Primary cutaneous follicle center cell lymphoma</td>
<td>2</td>
</tr>
<tr>
<td>EBV associated B cell proliferative disease</td>
<td>1</td>
</tr>
</tbody>
</table>