ANTIBODY PURIFICATION BY CATION EXCHANGE CHROMATOGRAPHY

Applicant: Genentech, Inc., South San Francisco, CA (US)

Inventors: Benedicte Andree Lebreton, San Francisco, CA (US); Deborah Ann O’Connor, San Carlos, CA (US); Aurelia Safta, Walnut Creek, CA (US); Mandakini Sharma, Sunnyvale, CA (US)

Assignee: Genentech, Inc., South San Francisco, CA (US)

Related U.S. Application Data

Continuation of application No. 14/531,880, filed on Nov. 3, 2014, now Pat. No. 9,896,478, which is a continuation of application No. 12/260,623, filed on Oct. 29, 2008, now abandoned.

Publication Classification

Int. Cl.
- C07K 1/18 (2006.01)
- C07K 1/36 (2006.01)
- C07K 16/22 (2006.01)
- C07K 16/06 (2006.01)
- C07K 16/28 (2006.01)

CPC C07K 1/18 (2013.01); C07K 2217/24 (2013.01); C07K 1/36 (2013.01); C07K 16/2887 (2013.01); C07K 16/22 (2013.01); C07K 16/065 (2013.01)

ABSTRACT

A method for purifying an antibody by cation exchange chromatography is described in which a high pH wash step is used to remove of contaminants prior to eluting the desired antibody using an elution buffer with increased conductivity.
<table>
<thead>
<tr>
<th></th>
<th>FR1</th>
<th></th>
<th></th>
<th>10</th>
<th>15</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ser</td>
<td>Gln Gly Val Leu Gln Gln Pro Gly Ala Glu Leu Val Lys Pro Gly Ala</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>20</td>
<td>25</td>
<td>30</td>
<td>31</td>
<td>35</td>
</tr>
<tr>
<td>Val Lys Met Ser Cys Lys Ala Ser Gly Tyr Thr Phe Thr Ser Tyr Asn Met His Trp</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>40</td>
<td>45</td>
<td>49</td>
<td>50</td>
<td>52</td>
</tr>
<tr>
<td>Val Lys Gin Thr Pro Gly Arg Gly Leu Glu Trp Ile Gly Ala Ile Tyr Pro Gly Asn</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>55</td>
<td>60</td>
<td>65</td>
<td>66</td>
<td>70</td>
</tr>
<tr>
<td>Gly Asp Thr Ser Tyr Asn Gin Lys Phe Lys Gly Lys Ala Thr Leu Thr Ala Asp Lys</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>75</td>
<td>80</td>
<td>82</td>
<td>82A</td>
<td>82B</td>
</tr>
<tr>
<td>Ser Ser Ser Thr Ala Tyr Met Gin Leu Ser Ser Leu Thr Ser Glu Asp Ser Ala Val</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>90</td>
<td>94</td>
<td>95</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tyr Tyr Cys Ala Arg Ser Thr Tyr Tyr Gly Asp Thr Tyr Phe Asn Thr Trp Gly</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>105</td>
<td>110</td>
<td>113</td>
<td>114</td>
<td></td>
</tr>
<tr>
<td>Ala Gly Thr Thr Val Thr Val Val Ser Ala Ala Ser Thr Lys Gly Pro Ser Val Phe Pro</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Human Gamma 1 Constant**

<p>| | | | | | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly Thr Ala Ala Leu Gly Cys Leu Val</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>130</td>
<td>133</td>
<td></td>
<td>140</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser Thr Ser Gly Ala Glu Ala Leu Thr Ser</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>150</td>
<td>154</td>
<td>156</td>
<td>157</td>
<td>162</td>
<td></td>
</tr>
<tr>
<td>Gly Val His Thr Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>169</td>
<td>171</td>
<td></td>
<td>180</td>
<td>182</td>
<td></td>
</tr>
<tr>
<td>Val Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gin Thr Tyr Ile Cys Asn Val Asn</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>190</td>
<td>200</td>
<td>203</td>
<td>205</td>
<td>210</td>
<td></td>
</tr>
<tr>
<td>His Lys Pro Ser Asn Thr Lys Val Asp Lys Lys Ala Glu Pro Lys Ser Cys Asp Lys</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>220</td>
<td>222</td>
<td>225</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Thr His Thr Cys Pro Pro Cys Pro Pro Ala Pro Glu Leu Leu Gly Gly Pro Ser Val Phe</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Human Gamma 1 Constant – continued

280 290 291 292
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val

295 296 299 300 310 314 317
Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser

320 330
Thr Tyr Arg Val Val Ser Val Leu Thr Val Val Leu His Glu Asp Trp Leu Asn Gly Lys

340 350 355
Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile

357 360 361 363 370
Ser Lys Ala Lys Gly Gin Pro Arg Glu Pro Gin Val Tyr Thr Leu Pro Pro Ser Arg

378 381 390
Asp Glu Leu Thr Lys Asn Gin Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro

408 410 413
Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gin Gin Pro Glu Asn Gin Tyr Lys Thr

420
Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val

428 430 433 440
Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val

448

450 460
Asp Lys Ser Arg Trp Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin Gin
Fig. 1B - Rituximab Light Chain

<table>
<thead>
<tr>
<th>7</th>
<th>8</th>
<th>9</th>
<th>10</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gln</td>
<td>Ile</td>
<td>Val</td>
<td>Leu Ser Glu Ser Pro Ala Ile Leu Ser Ala Ser</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>20</th>
<th>22</th>
<th>24</th>
<th>27</th>
<th>29</th>
<th>30</th>
<th>34</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pro Gly Glu Lys Val Thr Met Thr Cys Arg Ala Ser Ser Ser Val Ser Tyr Ile His</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>35</th>
<th>40</th>
<th>45</th>
<th>49</th>
<th>50</th>
</tr>
</thead>
<tbody>
<tr>
<td>CDR1 Thr Phe Gln Gln Lys Pro Gly Ser Ser Pro Lys Pro Thr Ile Tyr Ala Thr Ser Asn</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>55</th>
<th>56</th>
<th>57</th>
<th>60</th>
<th>63</th>
<th>70</th>
</tr>
</thead>
<tbody>
<tr>
<td>FR3 Len Ala Ser Gln Val Pro Val Arg Phe Ser Gly Ser Gly Ser Gly Thr Ser Tyr Ser</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>75</th>
<th>80</th>
<th>85</th>
<th>90</th>
<th>99</th>
<th>100</th>
</tr>
</thead>
<tbody>
<tr>
<td>CDR3 Thr Ser Asn Pro Gln Ala Ala Gln Ala Ala Thr Tyr Tyr Cys Gln Gln Trp</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>95</th>
<th>97</th>
<th>98</th>
<th>100</th>
<th>105</th>
<th>107</th>
<th>108</th>
<th>110</th>
</tr>
</thead>
<tbody>
<tr>
<td>FR4 Thr Ser Asn Pro Pro Thr Phe Gly Gln Gly Gln Thr Lys Gln Ala Ile Lys Arg Thr Val</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>120</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Gln Leu Lys Ser Gln Thr</td>
</tr>
</tbody>
</table>

**Human Kappa Constant**

<table>
<thead>
<tr>
<th>130</th>
<th>140</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Glu Trp</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>150</th>
<th>160</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gin Glu Ser Val Thr Glu Gin Asp</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>170</th>
<th>180</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>190</th>
<th>200</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glu Lys His Lys Val Tyr Ala Cys Gln Val Thr His Gin Gly Leu Ser Ser Ser Pro Val</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>210</th>
<th>214</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thr Lys Ser Phe Asn Arg Gly Glu Gln Thr</td>
<td></td>
</tr>
</tbody>
</table>

**Amino Acid # (kabat)**
Fig. 2A - Bevacizumab Heavy Chain

EVQLVESGGGLVQPGGLRLSCLAAASGTYFTNYGMNWVRQAPGK
GLEWVGWINTYTGEPTYAADFKRRRTFSLDTSKAYLQMNSLRA
EDTAVYYCAYKYPHYYGSSHWHYFDVGQTLVTVSSASTKGP
SFPLAPSSKSTSGGGTAAGLCVQKDVFPEPVTSVSNZGALTSGVH
TFPAVLQSSGLYSLSSVVTQPVSSLGLTQTYICNVTNIKPSNTKVDDKK
VEPKSCDKTHTCTCPCCCAPAILLLGQVSLFEPFPPKDPKETLMISRTPEVT
VVVDVSHEDEPEVKFNWYVWGVTVHINAKTTPREEQYNSTYRVS
VLTTLHQQDWNQNYKCKVSNKALPAPIEKTKQAKGQPREPQV
YLTIPPSREEMTNQAVSLTCVLKGFYPSDAVEWESNGQPENNYKT
TPPVLQSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALIINHT
QKSSLNLSPGK

Fig. 2B - Bevacizumab Light Chain

DIQMTQSPSSLSASYGDRVTITCSASQDISNYLNWYQQKPGKAPK
VLIYFTSSLGHSVPSPGSQGSDTFTLTISLQPEDFATYYCQQYS
TYPWTFGQGKTVEIKRTVAAPSIFIFPSDEQLKSGTASVCLLN
NFYPREAVQWVKDVNLGQSNQEVSWTEQDSKASTYSSSSLLTLS
KADYEKHKVYACEVTHQGLSSPVKSFNRGEC
FIG. 3 – Host Cell Proteins Removal for Rituximab Processes
ANTIBODY PURIFICATION BY CATION EXCHANGE CHROMATOGRAPHY

RELATED APPLICATIONS

[0001] This application claims the benefit of U.S. Provisional Patent Application No. 60/983,825, filed 30 Oct. 2007, the disclosure of which is hereby incorporated by reference in its entirety for all purposes.

BACKGROUND OF THE INVENTION

Field of the Invention

[0002] This invention relates generally to protein purification. In particular, the invention relates to a method for purifying antibody from a composition comprising the antibody and at least one contaminant using cation exchange chromatography, wherein a high pH wash step is used to remove contaminants prior to eluting the desired antibody using an elution buffer with increased conductivity.

Description of the Related Art

[0003] The large-scale, economic purification of proteins is an increasingly important problem for the biotechnology industry. Generally, proteins are produced by cell culture, using either eukaryotic or prokaryotic cell lines engineered to produce the protein of interest by insertion of a recombinant plasmid containing the gene for that protein. Since the cells typically used are living organisms, they must be fed with a complex growth medium, containing sugars, amino acids, and growth factors, usually supplied from preparations of animal serum. Separation of the desired protein from the mixture of compounds fed to the cells and from the by-products of the cells themselves to a purity sufficient for use as a human therapeutic poses a formidable challenge.

[0004] Procedures for purification of proteins from cell debris initially depend on the site of expression of the protein. Some protein can be secreted directly from the cell into the surrounding growth media; others are made intracellularly. For the latter proteins, the first step of a purification process involves lysis of the cell, which can be done by a variety of methods, including mechanical shear, osmotic shock, or enzymatic treatments. Such disruption releases the entire contents of the cell into the homogenate, and in addition produces subcellular fragments that are difficult to remove due to their small size. These are generally removed by differential centrifugation or by filtration. The same problem arises, although on a smaller scale, with directly secreted proteins due to the natural death of cells and release of intracellular host cell proteins in the course of the protein production run.

[0005] Once a clarified solution containing the protein of interest has been obtained, its separation from the other proteins produced by the cell is usually attempted using a combination of different chromatography techniques. These techniques separate mixtures of proteins on the basis of their charge, degree of hydrophobicity, or size. Several different chromatography resins are available for each of these techniques, allowing accurate tailoring of the purification scheme to the particular protein involved. The essence of each of these separation methods is that proteins can be caused either to move at different rates down a long column, achieving a physical separation that increases as they pass further down the column, or to adhere selectively to the separation medium, being then differentially eluted by different solvents. In some cases, the desired protein is separated from impurities when the impurities specifically adhere to the column, and the protein of interest does not, that is, the protein of interest is present in the “flow-through”.

[0006] Ion exchange chromatography is a chromatographic technique that is commonly used for the purification of proteins. In ion exchange chromatography, charged patches on the surface of the solute are attracted by opposite charges attached to a chromatography matrix, provided the ionic strength of the surrounding buffer is low. Elution is generally achieved by increasing the ionic strength (i.e., conductivity) of the buffer to compete with the solute for the charged sites of the ion exchange matrix. Changing the pH and thereby altering the charge of the solute is another way to achieve elution of the solute. The change in conductivity of pH may be gradual (gradient elution) or stepwise (step elution). In the past, these changes have been progressive, i.e., the pH or conductivity is increased or decreased in a single direction

[0007] U.S. Pat. Nos. 6,339,142, 6,417,355, 6,489,447, and 7,074,404 (Basey et al.) describe ion exchange chromatography for purifying polypeptides. U.S. Pat. Nos. 6,127,526, 6,333,398, and 6,797,814 (Blank, G.) describe purifying proteins, such as anti-HER2 antibodies, by Protein A chromatography. Methods for purifying proteins, such as antibodies, by ion exchange chromatography are described in U.S. Application Publication No. 2004/0082047.

SUMMARY OF THE INVENTION

[0009] The invention herein concerns an improved method for cation exchange chromatography of antibodies in which a high pH wash step is used to remove contaminants prior to eluting the desired antibody product. The process results, amongst other things, in improved removal of Chinese Hamster Ovary Proteins (CHOP) contaminants.

[0010] According to a first aspect, the invention provides a method for purifying an antibody from a composition comprising the antibody and at least one contaminant, which method comprises the sequential steps of:

[0011] (a) loading the composition onto a cation exchange material wherein the composition is at a first pH;

[0012] (b) washing the cation exchange material with a first wash buffer at a pH which is greater than that of the composition in (a), wherein the pH of the first wash buffer is from about 6.8 to about 9.0;

[0013] (c) washing the cation exchange material, with a second wash buffer at a pH which is less than that of the first wash buffer; and

[0014] (d) eluting the antibody from the cation exchange material with an elution buffer at a conductivity which is substantially greater than that of the second wash buffer.

[0015] Preferably the antibody binds human CP20, such as rituximab, or binds human vascular endothelial growth factor (VEGF), such as bevacizumab.

[0016] According to one preferred embodiment, the invention concerns a method for purifying an antibody that binds human CD20 from a composition comprising the antibody and one or more contaminants selected from the group consisting of Chinese Hamster Ovary Proteins (CHOP), leached protein A, DNA, and aggregated CD20 antibody, which method comprises the sequential steps of:

[0017] (a) loading the composition onto a cation exchange material wherein the composition is at a pH from about 4.0 to about 6.0;

[0018] (b) washing the cation exchange material with a first wash buffer at a pH from about 6.8 to about 9.0;

[0019] (c) washing the cation exchange material with a second wash buffer at a pH from about 5.0 to about 6.0; and

[0020] (d) eluting the antibody from the cation exchange material using an elution buffer with a pH from about 5.0 to about 6.0 and a conductivity from about 10 to about 100 mS/cm. Preferably the CD20 antibody is rituximab.

[0021] According to another preferred embodiment, the invention relates to a method for purifying an antibody that binds human vascular endothelial growth factor (VEGF) from a composition comprising the antibody and one or more contaminants selected from the group consisting of a cell culture media component, garamycin, Chinese Hamster Ovary Proteins (CHOP), DNA, viral contaminant, and aggregated VEGF antibody, which method comprises the sequential steps of:

[0022] (a) loading the composition onto a cation exchange material wherein the composition is at a pH from about 4.0 to about 6.0;

[0023] (b) washing the cation exchange material with a first wash buffer at a pH from about 6.8 to about 8.0;

[0024] (c) washing the cation exchange material with a second wash buffer at a pH from about 5.0 to about 6.0; and

[0025] (d) eluting the antibody from the cation exchange material using an elution buffer with a pH from about 5.0 to about 6.0 and a conductivity from about 10 to about 100 mS/cm. Preferably, the VEGF antibody is bevacizumab.

[0026] The invention also concerns a composition comprising rituximab in a buffer comprising about 25 mM HEPES, at a pH of about 7.8.

[0027] In addition, the invention provides a composition comprising bevacizumab in a buffer comprising about 25 mM MOPS at a pH of about 7.0.

BRIEF DESCRIPTION OF THE DRAWINGS

[0028] FIGS. 1A and 1B provide the amino acid sequences of the heavy chain (SEQ ID No. 1) and light chain (SEQ ID No. 2) of rituximab antibody. Each of the framework regions (FR1-4) and each of the CDR regions (CDR1-3) in each variable region are identified, as are the human gamma 1 heavy chain constant sequence and human kappa light chain constant sequence. The variable heavy (VH) region is in SEQ ID No. 3. The variable light (VL) region is in SEQ ID No. 4. The sequence identifiers for the CDRs are: CDR H1 (SEQ ID No. 5), CDR H2 (SEQ ID No. 6), CDR H3 (SEQ ID No. 7), CDR L1 (SEQ ID No. 8), CDR L2 (SEQ ID No. 9), and CDR L3 (SEQ ID No. 10).

[0029] FIGS. 2A and 2B provide the amino acid sequences of the heavy chain (SEQ ID No. 11) and light chain (SEQ ID No. 12) of bevacizumab antibody. The end of each variable region is indicated with ||. The variable heavy (VH) region is in SEQ ID No. 13. The variable light (VL) region is in SEQ ID No. 14. Each of the three CDRs in each variable region is underlined. The sequence identifiers for the CDRs are: CDR H1 (SEQ ID No. 15), CDR H2 (SEQ ID No. 16), CDR H3 (SEQ ID No. 17), CDR L1 (SEQ ID No. 18), CDR L2 (SEQ ID No. 19), and CDR L3 (SEQ ID No. 20).

[0030] FIG. 3 provides a side-by-side comparison of host cell proteins removal by the cation exchange chromatography process of the improved rituximab processed compared to the original process. Superior CHOP removal was achieved with the new process.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

Definitions:

[0031] Herein, numerical ranges or amounts prefixed by the term “about” expressly include the exact range or exact numerical amount.

[0032] The “composition” to be purified herein comprises the antibody of interest and one or more contaminants. The composition may be “partially purified” (i.e. having been subjected to one or more purification steps) or may be obtained directly from a host cell or organism producing the antibody (e.g. the composition may comprise harvested cell culture fluid).
As used herein, “polypeptide” refers generally to peptides and proteins having more than about ten amino acids. Preferably, the polypeptide is a mammalian protein, examples of which include: renin; a growth hormone, including human growth hormone and bovine growth hormone; growth hormone releasing factor; parathyroid hormone; thyroid stimulating hormone; lipoproteins; alpha-1-antitrypsin; insulin A-chain; insulin B-chain; proinsulin; follicle stimulating hormone; calcitonin; luteinizing hormone; glucagon; clotting factors such as factor VIIc; factor IX, tissue factor; and von Willebrand factor; anti-clotting factors such as Protein C; atrial natriuretic factor; lung surfactant; a plasminogen activator, such as urokinase or human urine or tissue-type plasminogen activator (t-PA); bombesin; thrombin; hemopoietic growth factor; tumor necrosis factor-alpha and -beta; enkephalins; RANTES (regulated on activation normally T-cell expressed and secreted); human macrophage inflammatory protein (MIP-1-alpha); a serum albumin such as human serum albumin; Muellerian-inhibiting substance; relaxin A-chain; relaxin B-chain; preprorelaxin; mouse gonadotropin-associated peptide; a microbial protein, such as beta-lactamase; DNase; IgE; a cytotoxic T-lymphocyte associated antigen (CTLA), such as CTLA-4; inhibin; activin; vascular endothelial growth factor (VEGF); receptors for hormones or growth factors; Protein A or D; rheumatoid factors; a neutrophilic factor such as bone-derived neurotrophic factor (BDNF), neurotrophin-3, -4, -5, or -6 (NT-3, NT-4, NT-5, or NT-6), or a nerve growth factor such as NGF-B; platelet-derived growth factor (PDGF); fibroblast growth factor such as aFGF and bFGF; epidermal growth factor (EGF); transforming growth factor (TGF) such as TGF-alpha and TGF-beta, including TGF-β1, TGF-β2, TGF-β3, TGF-β4, or TGF-β5; insulin-like growth factor I and II (IGF-I and IGF-II); des(1-3)-IGF-1 (brain IGF-1), insulin-like growth factor binding proteins (IGFBPs); CD proteins such as CD3, CD4, CD8, CD19 and CD20; erythropoietin; osteoinductive factors; immunotoxins; a bone morphogenetic protein (BMP); an interferon such as interferon-alpha, -beta, and -gamma; colony stimulating factors (CSFs), e.g., M-CSF, GM-CSF, and G-CSF; interleukins (ILs), e.g., IL-1 to IL-10; superoxide dismutase; T-cell receptors; surface membrane proteins; decay accelerating factor; viral antigen such as, for example, a portion of the AIDS envelope; transport proteins; home receptors; addressins; regulatory proteins; integrins such as CD11a, CD11b, CD11c, CD18, an ICAM, VLA-4 and VCAM; a tumor associated antigen such as HER2, HER3 or HER4 receptor; and fragments and/or variants of any of the above-listed polypeptides as well as antibodies, including antibody fragments, binding to any of the above-listed polypeptides. A preferred polypeptide is an intact antibody or an antibody fragment that binds to human CD20, for example, rituximab; or an intact antibody or an antibody fragment that binds to human vascular endothelial growth factor (VEGF), for example bevacizumab.

The phrase “cation exchange material” refers to a solid phase that is negatively charged and has free cations for exchange with cations in an aqueous solution passed over or through the solid phase. The charge may be provided by attaching one or more charged ligands to the solid phase, e.g., by covalent linking. Alternatively, or in addition, the charge may be an inherent property of the solid phase (e.g., as is the case for silica, which has an overall negative charge). Commercially available cation exchange materials include carboxy-methyl-cellulose, BAKERBOND ABX™, sulfopropyl (SP) immobilized on agarose (e.g., SP-SEP-HAROSE FAST FLOW™, SP-SEP-HAROSE FAST FLOW XL™ or SP-SEP-HAROSE HIGH PERFORMANCE™, from GE Healthcare), CAPTO S™ (GE Healthcare), FACTOGEL-SO3™, FACTOGEL-SE HICAP™, and FRACTO-PREP™ (EMD Merck), sulphonyl immobilized on agarose (e.g., S-SEP-HAROSE FAST FLOW™ from GE Health-care), and SUPER SP™ (Tosoh Biosciences). A preferred cation exchange material herein comprises cross-linked poly(styrene-divinylbenzene) flow-through particles (solid phase) coated with a polyhydroxylated polymer functionalized with sulfopropyl groups (for example, POROS 50 HS® chromatography resin).

By “solid phase” is meant non-aqueous matrix to which one or more charged ligands can adhere. The solid phase may be a purification column (including, without limitation, expanded bed and packed bed columns), a discontinuous phase of discrete particles, a membrane, or filter etc. Examples of materials for forming the solid phase include polysaccharides such as agarose and cellulose) and other mechanically stable matrices such as silica (e.g. controlled pore glass), poly(styrene-divinylbenzene), polycarylamide, ceramic particles and derivatives of any of the above.

The term “load” herein refers to the composition loaded onto the cation exchange material. Preferably, the cation exchange material is equilibrated with an equilibration buffer prior to loading the composition which is to be purified.

A “buffer” is a solution that resists changes in pH by the action of its acid-base conjugate components. Various buffers which can be employed depending, for example, on the desired pH of the buffer are described in Buffers: A guide for the Preparation and Use of Buffers in Biological Systems, Guefroy, D., Ed. Calbiochem Corporation (1975).

An “equilibration buffer” is a buffer that is used to equilibrate the cation exchange material, prior to loading the composition comprising the antibody of interest and one or more contaminants onto the cation exchange material. Preferably the pH of the equilibration buffer herein is in the range from about 5.0 to about 6.0, preferably about 5.5. Preferably, the conductivity of the equilibration buffer herein is in the range from about 1 to about 8 mS/cm, preferably from about 4 to about 8 mS/cm, and most preferably from about 5 to about 8 mS/cm. Optionally, the equilibration buffer comprises a salt, such as NaCl, for example, in an amount from about 40 mM to about 80 mM, preferably about 60 mM NaCl.

The term “wash buffer” is used herein to refer to the buffer that is passed over the cation exchange material following loading of a composition and prior to elution of the protein of interest. The wash buffer may serve to remove one or more contaminants from the cation exchange material, without substantial elution of the desired antibody.
product. According to the preferred embodiment of the invention wherein a "first wash buffer" and a "second wash buffer" are used.

[0041] Herein, the expression "first wash buffer" refers to a wash buffer having a pH increased relative to the pH of the load and/or equilibration buffer. The first wash buffer may be used herein to elute one or more contaminants from the cation exchange material, without substantially eluting the antibody product of interest therefrom. The term "first" should not be interpreted as excluding the use of one or more additional wash or other buffers between the load and the first wash buffer. Preferably the pH of the first wash buffer herein is in the range from about 6.8 to about 9.0, preferably from about 7.0 to about 8.0, and most preferably pH about 7.0 or pH about 7.8. Preferably, the conductivity of the first wash buffer herein is in the range from about 0.01 to about 5 mS/cm, preferably from about 0.1 to about 3 mS/cm, and most preferably from about 0.2 to about 2 mS/cm. Optionally, the first wash buffer is substantially free of a salt (such as NaCl) therein.

[0042] The expression "second wash buffer" for the purposes of this application refers to a wash buffer used after the first wash buffer to prepare the cation exchange material for elution of the antibody of interest. The term "second" should not be interpreted as excluding the use of one or more additional wash or other buffers between the first wash buffer and second wash buffer. Preferably the pH of the second wash buffer herein is in the range from about 5.0 to about 6.0, preferably about 5.5, and most preferably pH 5.5. Preferably, the conductivity of the second wash buffer herein is in the range from about 0.01 to about 5 mS/cm, preferably about 0.1 to about 3 mS/cm, and most preferably from about 0.5 to about 3.0 mS/cm.

[0043] "Elution buffer" is used to elute the antibody of interest from the solid phase. Herein, the elution buffer has a substantially increased conductivity relative to that of the second wash buffer, such that the desired antibody product is eluted from the cation exchange material. Preferably, the conductivity of the elution buffer is substantially greater than that of the load and of each of the preceding buffers, namely of the equilibration buffer, first wash buffer, and second wash buffer. By "substantially greater" conductivity is meant, for example, that the buffer has a conductivity which is at least 2, 3, 4, 5 or 6 conductivity units (mS/cm) greater than that of the composition or buffer to which it is being compared. In one embodiment, the pH of the elution buffer is substantially the same as that of the equilibration and/or second wash buffer. Preferably the pH of the elution buffer herein is in the range from about 5.0 to about 6.0, preferably about 5.5, and most preferably pH 5.5. Preferably, the conductivity of the elution buffer herein is in the range from about 10 mS/cm to about 100 mS/cm, preferably from about 12 mS/cm to about 30 mS/cm, and most preferably from about 12 to about 20 mS/cm. Increased conductivity may be achieved by the addition of a salt, such as sodium chloride, sodium acetate, potassium chloride to the elution buffer. Preferably, the elution buffer comprises from about 100 to about 300 mM NaCl, preferably from about 150 mM to about 200 mM NaCl, for example about 175 mM NaCl or about 160 mM NaCl.

[0044] A "regeneration buffer" may be used to regenerate the cation exchange material such that it can be re-used. The regeneration buffer has a conductivity and/or pH as required to remove substantially all contaminants and the antibody of interest from the cation exchange material.

[0045] The term "conductivity" refers to the ability of an aqueous solution to conduct an electric current between two electrodes. In solution, the current flows by ion transport. Therefore, with an increasing amount of ions present in the aqueous solution, the solution will have a higher conductivity. The basic unit of measure for conductivity is the Siemens (or mho), mho (mS/cm), and can be measured using a conductivity meter, such as various models of Orion conductivity meters. Since electrolytic conductivity is the capacity of ions in a solution to carry electrical current, the conductivity of a solution may be altered by changing the concentration of ions therein. For example, the concentration of a buffering agent and/or the concentration of a salt (e.g. sodium chloride, sodium acetate, or potassium chloride) in the solution may be altered in order to achieve the desired conductivity. Preferably, the salt concentration of the various buffers is modified to achieve the desired conductivity.

[0046] By "purifying" an antibody from a composition comprising the antibody and one or more contaminants is meant increasing the degree of purity of the antibody in the composition by removing (completely or partially) at least one contaminant from the composition. A "purification step" may be part of an overall purification process resulting in a "homogeneous" composition. "Homogeneous" is used herein to refer to a composition comprising at least about 70% by weight of the antibody of interest, based on total weight of the composition, preferably at least about 80% by weight, more preferably at least about 90% by weight, even more preferably at least about 95% by weight.

[0047] By "binding" a molecule to a cation exchange material is meant exposing the molecule to the cation exchange material under appropriate conditions (pH and/or conductivity) such that the molecule is reversibly immobilized in or on the cation exchange material by virtue of ionic interactions between the molecule and a charged group or charged groups of the cation exchange material.

[0048] By "washing" the cation exchange material is meant passing an appropriate buffer through or over the cation exchange material.

[0049] By "eluting" a molecule (e.g., antibody or contaminant) from a cation exchange material is meant to remove the molecule therefrom.

[0050] In preferred embodiments of the invention, the antibody to be purified herein is a recombinant antibody. A "recombinant antibody" is one which has been produced in a host cell which has been transformed or transfected with nucleic acid encoding the antibody, or produces the antibody as a result of homologous recombination. "Transformation" and "transfection" are used interchangeably to refer to the process of introducing nucleic acid into a cell. Following transformation or transfection, the nucleic acid may integrate into the host cell genome, or may exist as an extrachromosomal element. The "host cell" includes a cell in in vitro cell culture as well as a cell within a host animal. Methods for recombinant production of polypeptides are described in U.S. Pat. No. 5,534,615, expressly incorporated herein by reference, for example.

[0051] A "variant" or "amino acid sequence variant" of a starting polypeptide is a polypeptide that comprises an amino acid sequence different from that of the starting polypeptide. Generally, a variant will possess at least 80%
sequence identity, preferably at least 90% sequence identity, more preferably at least 95% sequence identity, and most preferably at least 98% sequence identity with the native polypeptide. Percentage sequence identity is determined, for example, by the Fitch et al., *Proc. Natl. Acad. Sci. USA* 80:1382-1386 (1983), version of the algorithm described by Needleman et al., *J. Mol. Biol.* 48:443-453 (1970), after aligning the sequences to provide for maximum homology. Amino acid sequence variants of a polypeptide may be prepared by introducing appropriate nucleotide changes into DNA encoding the polypeptide, or by peptide synthesis. Such variants include, for example, deletions from, and/or insertions into and/or substitutions of, residues within the amino acid sequence of the polypeptide of interest. Any combination of deletion, insertion, and substitution is made to arrive at the final construct, provided that the final construct possesses the desired characteristics. The amino acid changes also may alter post-translational processing of the polypeptide, such as by changing the number or position of glycosylation sites. Other post-translational modifications include hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl, threonyl or tyrosyl residues, methylation of the α-amino groups of lysine arginine and histidine side chains (T. E. Creighton, Proteins: Structure and Molecular Properties, W.H. Freeman & Co., San Francisco, pp. 79-86 (1983)). Methods for generating amino acid sequence variants of polypeptides are described in U.S. Pat. No. 5,534,615, expressly incorporated herein by reference, for example.

The term “antibody” is used in the broadest sense and specifically covers monoclonal antibodies (including full length monoclonal antibodies), polyclonal antibodies, multispecific antibodies (e.g., bispecific antibodies), and antibody fragments so long as they exhibit the desired binding specificity.

The antibody herein is directed against an “anti-gen” of interest. Preferably, the antigen is a biologically important polypeptide and administration of the antibody to a mammal suffering from a disease or disorder can result in a therapeutic benefit in that mammal. However, antibodies directed against non-polypeptide antigens (such as tumor-associated glycolipid antigens; see U.S. Pat. No. 5,091,178) are also contemplated. Where the antigen is a polypeptide, it may be a transmembrane molecule (e.g., receptor) or ligand such as a growth factor. Exemplary antigens include those polypeptides discussed above. Preferred molecular targets for antibodies encompassed by the present invention include CD polypeptides such as CD3, CD4, CD8, CD19, CD20 and CD34; members of the HER receptor family such as the EGF receptor (HER1), HER2, HER3 or HER4 receptor; cell adhesion molecules such as LFA-1, Mac1, p150,95, VLA-4, ICAM-1, VCAM and avb3 integrin including either a or b subunits thereof (e.g. anti-CD11a, anti-CD18 or anti-CD11b antibodies); growth factors such as VEGF; IgE; blood group antigens; ifk2/fk3 receptor; obesity (OB) receptor; mpl receptor; CTLA-4; polypeptide C etc. Soluble antigens or fragments thereof, optionally conjugated to other molecules, can be used as immunogens for generating antibodies. For transmembrane molecules, such as receptors, fragments of these (e.g. the extracellular domain of a receptor) can be used as the immunogen. Alternatively, cells expressing the transmembrane molecule can be used as the immunogen. Such cells can be derived from a natural source (e.g., cancer cell lines) or may be cells which have been transformed by recombinant techniques to express the transmembrane molecule.

Examples of antibodies to be purified herein include, but are not limited to: HER2 antibodies including trastuzumab (HERCEPTIN®) (Carter et al., *Proc. Natl. Acad. Sci. USA* 89:4285-4289 (1989), U.S. Pat. No. 5,725,866 and pertuzumab (OMNITARG™) (WO01/00245); CD20 antibodies (see below); IL-8 antibodies (St John et al., *Chest* 103:932 (1993), and International Publication No. WO 95/23865); VEGF or VEGF receptor antibodies including humanized and/or affinity matured VEGF antibodies such as the humanized VEGF antibody huA4.6.1 bevaciuzumab (AVASTIN®) and ranibizumab (LUCENTIS®) (Kim et al., *Growth Factors*, 7:53-64 (1992), International Publication No. WO 96/30086, and WO 98/45331, published Oct. 15, 1998); PSMA antibodies (WO99/40369); CD11a antibodies including eflazuzumab (RAPTIVA®) (U.S. Pat. No. 5,622,700; WO 98/23761, Steppe et al., Transplantation Int'l 4:3-7 (1991), and Hourmant et al., Transplantation 58:377-380 (1994)); antibodies that bind IgE such as omalizumab (XOLAIR®) (Presta et al., *J. Immunol.* 151: 2523-2632 (1993), and International Publication No. WO 95/19181; U.S. Pat. No. 5,714,338, issued Feb. 3, 1998 and U.S. Pat. No. 5,001,313, issued Feb. 25, 1992, WO 93/04173 published Mar. 4, 1993, or International Application No. PCT/US98/13410 filed Jun. 30, 1998, U.S. Pat. No. 5,714,338); CD18 antibodies (U.S. Pat. No. 5,622,700, issued Apr. 22, 1997, or as in WO 97/26912, published Jul. 31, 1997); Apo-2 receptor antibody antibodies (WO 98/51793 published Nov. 19, 1998); Tissue Factor (TF) antibodies (European Patent No. 0 420 937 B1 granted Nov. 9, 1994); αv-α, integrin antibodies (WO 98/06248 published Feb. 19, 1998); EGF receptor antibodies (e.g., chimerized or humanized 225 antibody, cetuximab, ERBUTIX® as in WO 96/40210 published Dec. 19, 1996); CD3 antibodies such as OKT3 (U.S. Pat. No. 4,515,893 issued May 7, 1985); CD25 or Tac antibodies such as CH1-621 (SIMULECT®) and ZENA-PAX® (See U.S. Pat. No. 5,693,762 issued Dec. 2, 1997); CD4 antibodies such as the CM-7412 antibody (Choy et al. *Arthritis Rheum.* 39(1):52-56 (1996)); CD52 antibodies such as CAMPATH-III (ILEX/Belex) (Reichmann et al. *Nature* 323:323-337 (1986); Fe receptor antibodies such as the M22 antibody directed against FeR1 as in Graziano et al. *J. Immunol.* 155(10):4996-5002 (1995); carcinoembryonic antigen (CEA) antibodies such as iMN-14 (Sharkey et al. *Cancer Res.* 55(23Suppl): 5935s-5945s (1995)); antibodies directed against breast epithelial cells including hu1bRl-3, hu-Mc 3 and CHL6 (Cerami et al. Cancer Res. 55(23): 5852s-5856s (1995)) and Richman et al. Cancer Res. 55(23 Supp): 5916a-5920s (1995)); antibodies that bind to colon carcinoma cells such as C242 (Litton et al. *Eur. J. Immunol.* 26(1):1-9 (1996)); CD38 antibodies, e.g. AT 13/5 (Ellis et al. *J. Immunol.* 155(2):925-937 (1995)); CD33 antibodies such as Hu M195 (Juricke et al. Cancer Res 55(23 Suppl):5908s-5910s (1995)) and CMA-676 or CDP771; EpCAM antibodies such as 17-1A (PANOREX®); GplblbIIa antibodies such as abeximab or c7E3 Fab (REOPOR®, RSV antibodies such as MEDI-493 (SYNAGIS®); CMV antibodies such as PROTOVIR®; HIV antibodies such as PRO542; hepatitis antibodies such as the hlep B antibody OSTAVIR®; CA 125 antibody OvaRel; idiotype CD3 epitope antibody BEC2; αvβ3 antibody (e.g. VITAXIN®, Medimmune); human renal cell carcinoma hormone 4 structure as ch-G250; ING-1; anti-human 17-1A antibody (3622W94); anti-lu-
man colorectal tumor antibody (A33); anti-human melanoma antibody R24 directed against GD3 ganglioside; anti-human squamous-cell carcinoma (SF-25); human leukocyte antigen (HLA) antibody such as Smart ID10 and the anti-HLA DR antibody Oncoyl (1-ym-1); CD37 antibody such as TRU 016 (Trubion); IL-21 antibody (Zymogenetics/Novo Nordisk); anti-B cell antibody (Impheron); B cell targeting MAb (Immunogen/Aventis); 1D90C3 (Morphyos/GPC); LymphoRad 131 (HGS); LY-1 antibody, such as LYM-1-1Y-90 (USC) or anti-Lym-1 Oncoly (USC/Peregrine); LIF 226 (Enhanced Lifesci.); BAFF antibody (e.g., WO 03/36358); BAFF receptor antibody (see e.g., WO 02/24090); BR3 antibody; Blys antibody such as belimumab; LYMHPoSTAT-BTM; ISF 154 (USCD/Roche/Tra- gen); gomilixima (Idex 152; Biogen Idec); IL-6 receptor antibody such as atiluzumab (ACETMRA™; Chugai/Roche); IL-15 receptor antibody such as HuMax-15-15 (Genmab/Angen); chemokine receptor antibody, such as a CCRL2 receptor (e.g., MLN1202; Millenium); anti-complement antibody, such as C5 antibody (e.g., ecuclizumab, 5G11.Alexion); oral infection of human immunoglobulin (e.g., IgPQ; Protein therapeutics); IL-12 receptor antibody such as ABT-874 (CAT/Abbott); Tenexilmax (BMS-224818; BMS); CD40 antibodies, including 2Z6 and humanized variants thereof (WO/00/ 75348) and TNX 100 (Chiron/Tanox); TNF-α antibodies including cA2 or infliximab (REMICADE©); CDP 571, MAK-195, adalimumab (HUMIRA™), pegylated TNF-α antibody fragment such as CDP-870 (Celltech), D2E7 (Knoll), anti-TNF-α polyclonal antibody (e.g., PassTNF; Verigen); CD22 antibodies such as L.1.2 or epratuzumab (LYMPHOCIDE©; Immunomedics), including adalimumab Y-90 and epratuzumab 1-131, Abiogen’s CD22 antibody (Abiogen, Italy), CMC 544 (Wythell/Celltech), tox (UT Southwestern), BL22 (NIH), ad LymopoScan Tc99 (Immunomedics). Preferably, the antibody that is purified herein is a naked, intact antibody which binds to human CD20, or a naked, intact antibody which binds to human VEGF.

[0055] The human “CD20” antigen, or “CD20,” is an about 35-kDa, non-glycosylated phosphoprotein found on the surface of greater than 90% of B cells from peripheral blood or lymphoid organs. CD20 is present on both normal B cells as well as malignant B cells, but is not expressed on stem cells. Other names for CD20 in the literature include “B-lymphocyte-restricted antigen” and “Bp35”. The CD20 antigen is described in Clark et al., Proc. Natl. Acad. Sci. (USA) 82:1766 (1985), for example.

[0056] A “CD20 antibody antagonist” herein is an antibody that, upon binding to CD20 on B cells, destroys or depletes B cells in a subject and/or interferes with one or more B-cell functions, e.g., by reducing or preventing a humoral response elicited by the B cell. The antibody antagonist preferably is able to deplete B cells (i.e., reduce circulating B-cell levels) in a subject treated therewith. Such depletion may be achieved by various mechanisms such as antibody-dependent cell-mediated cytotoxicity (ADCC) and/or complement-dependent cytotoxicity (CDC), inhibition of B-cell proliferation and/or induction of B-cell death (e.g., via apoptosis).

[0057] As used herein, “B cell depletion” refers to a reduction in B cell levels in an animal or human generally after drug or antibody treatment, as compared to the level before treatment. B cell depletion can be partial or complete. B cell levels are measurable using well known techniques such as those described in Reff et al., Blood 83: 435-445 (1994), or U.S. Pat. No. 5,736,137 (Anderson et al.). By way of example, a mammal (e.g. a normal primate) may be treated with various dosages of the antibody or immunoadhesin, and peripheral B-cell concentrations may be determined, e.g. by a FACS method that counts B cells.

[0058] Examples of CD20 antibodies include: “C2B8,” which is now called “rituximab” (“RITUXAN®”/”RITUXAN®”/”RITUXAN®”) (U.S. Pat. No. 5,736,137); the yttrium-[90]labeled 218 murine antibody designated “Y2B8” or “Ibritumomab Tuxetan” (ZEN- VALIN®) commercially available from IDEC Pharmaceuticals, Inc. (U.S. Pat. No. 5,736,137; 218 deposited with ATCC under accession no. HB11388 on Jun, 22, 1993); murine IgG2a “B1,” also called “Tositumomab,” optionally labelled with 131I to generate the “13Il-B1” or “iodule I13I tositumomumab” antibody (BEXXAR™) commercially available from Corixa (see, also, U.S. Pat. No. 5,595,721); murine monoclonal antibody “1F5” (Press et al. Blood 69(2):584-581 (1987) and variants thereof including “framework patched” or humanized 1F5 (WO 2003/002607, Leung, S.; ATCC deposit HB-96450); murine 2H7 and chimeric 2H7 antibody (U.S. Pat. No. 5,677,180); humanized 2H7 (WO 2004/056312, Lowman et al., and as set forth below); 2F2 (HuMax-CD20), a fully human, high-affinity antibody targeted at the CD20 molecule in the cell membrane of B-cells (Genmab, Denmark; see, for example, Glennie and van de Winkel, Drug Discovery Today 8: 503-510 (2003) and Cragg et al., Blood 101: 1045-1052 (2003); WO 2004/035607; US2004/0167319); the human monoclonal antibodies set forth in WO 2004/035607 and US2004/0167319 (Teeling et al.) the antibodies having complex N-glycoside-linked sugar chains bound to the Fe region described in US 2004/0093621 (Shitara et al.); monoclonal antibodies and antigen-binding fragments binding to CD20 (WO 2005/000901, Tedder et al.) such as HB20-3, HB20-4, HB20-5, and MB20-11; CD20 binding molecules such as the AME series of antibodies, e.g., AME 33 antibodies as set forth in WO 2004/103404 and US2005/ 0025764 (Watkins et al., Eli Lilly/Applied Molecular Evolution, AME); CD20 binding molecules such as those described in US 2005/0025764 (Watkins et al.); A20 antibody or variants thereof such as chimeric or humanized A20 antibody (cA20, hA20, respectively) or IMMU-106 (US 2003/0219453, Immunomedics); CD20-binding antibodies, including epitope-depleted Leu-16, 1H4, or 2B8, optionally conjugated with IL-2, as in US 2005/0069545A1 and US 2005/16969 (Carr et al.) bispecific antibody that binds CD22 and CD20, for example hL2:hA20 (WO2005/ 14618, Chang et al.); monoclonal antibodies L27, G28-2, 93-118, B-C1 or NU-B2 available from the International Leukocyte Typing Workshop (Valentine et al., In: Leukocyte Typing III (McMichael, Ed., p. 440, Oxford University Press (1987)); 11F4 (Haisma et al. Blood 92:184 (1998)); anti-CD20 aspartate E conjugate (Seattle Genetics); anti-CD20-IL2 (EMD/Biovation/City of Hope); anti-CD20 MAB (EpiCye); anti-CD20 antibody TRU 015 (Trubion). The preferred CD20 antibodies herein are chimeric, humanized, or human CD20 antibodies, more preferably rituximab, humanized 218, 212 (Hu-Max-CD20) human CD20 antibody (Genmab), and humanized A20 or IMMU-106 antibody (Immunomedics).

[0059] For the purposes herein, the terms “rituximab,” “RITUXAN®,” and “C2B8” herein refer to a recombinant chimeric antibody which binds to the human CD20 antigen.
as described in U.S. Pat. No. 5,736,137, Anderson et al. Such antibody preferably comprises a heavy chain comprising CDR H1 (SEQ ID No. 5), CDR H2 (SEQ ID No. 6), CDR H3 (SEQ ID No. 7), and a light chain, wherein the light chain preferably comprises CDR L1 (SEQ ID No. 8), CDR L2 (SEQ ID No. 9), and CDR L3 (SEQ ID No. 10); preferably the heavy chain comprises a variable heavy (VH) region comprising SEQ ID No. 3 and a variable light (VL) region comprising SEQ ID No. 4; and most preferably comprises a heavy chain comprising SEQ ID No. 1 (with or without a C-terminal lysine residue), and a light chain, wherein the light chain preferably comprises SEQ ID No. 2.

The terms expressly include variant forms such as described in Moorhouse et al. J. Pharm Biomed. Anal. 16:593-603 (1997).

[0060] The term “human VEGF” as used herein refers to the 165-amino acid human vascular endothelial cell growth factor, and related 121-, 189-, and 206-amino acid vascular endothelial cell growth factors, as described by Leung et al., Science 246:1306 (1989), and Houch et al., Mol. Endocrin. 5:1806 (1991) together with the naturally occurring allelic and processed forms of those growth factors.

[0061] The present invention provides anti-VEGF antagonistic antibodies which are capable of inhibiting one or more of the biological activities of VEGF, for example, its mitogenic or angiogenic activity. Antagonists of VEGF act by interfering with the binding of VEGF to a cellular receptor, by incapacitating or killing cells which have been activated by VEGF, or by interfering with vascular endothelial cell activation after VEGF binding to a cellular receptor. All such points of intervention by a VEGF antagonist shall be considered equivalent for purposes of this invention.

[0062] For the purposes herein, the terms “bevacizumab,” “AVASTIN®,” “F(ab)2,” and “ruMAb VEGF” herein refer to a recombinant humanized monoclonal antibody which binds human vascular endothelial growth factor (VEGF) antigen (ruMAb VEGF) as described in U.S. Pat. No. 7,169,901, Presta et al. Such antibody preferably comprises a heavy chain comprising CDR H1 (SEQ ID No. 15), CDR H2 (SEQ ID No. 16), CDR H3 (SEQ ID No. 17), and a light chain, wherein the light chain preferably comprises CDR L1 (SEQ ID No. 18), CDR L2 (SEQ ID No. 19), and CDR L3 (SEQ ID No. 20); most preferably the heavy chain comprises a variable heavy (VH) region comprising SEQ ID No. 3 and a variable light (VL) region comprising SEQ ID No. 4; and preferably comprises a heavy chain comprising SEQ ID No. 11 (with or without a C-terminal lysine residue), and a light chain, wherein the light chain preferably comprises SEQ ID No. 12. The terms expressly include variant forms that form during production of the recombinant antibody product.

[0063] The term “monoclonal antibody” as used herein refers to an antibody obtained from a population of substantially homogeneous antibodies, i.e., the individual antibodies comprising the population are identical except for possible naturally occurring mutations that may be present in minor amounts. Monoclonal antibodies are highly specific, being directed against a single antigenic site. Furthermore, in contrast to conventional (polyclonal) antibody preparations which typically include different antibodies directed against different determinants (epitopes), each monoclonal antibody is directed against a single determinant on the antigen. The modifier “monoclonal” indicates the character of the antibody as being obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method. For example, the monoclonal antibodies to be used in accordance with the present invention may be made by the hybridoma method first described by Kohler et al., Nature 256:495 (1975), or may be made by recombinant DNA methods (see, e.g., U.S. Pat. No. 4,816,567). In a further embodiment, “monoclonal antibodies” can be isolated from antibody phage libraries generated using the techniques described in McCafferty et al., Nature 348:552-554 (1990). Clarkson et al., Nature, 352:624-628 (1991) and Marks et al., J. Mol. Biol., 222:581-597 (1991) describe the isolation of murine and human antibodies, respectively, using phage libraries. Subsequent publications describe the production of high affinity (nM range) human antibodies by chain shuffling (Marks et al., BioTechnology, 10:779-783 (1992)), as well as combinatorial infection and in vivo recombination as a strategy for constructing very large phage libraries (Waterhouse et al., Nuc. Acids. Res., 21:2265-2266 (1993)). Thus, these techniques are viable alternatives to traditional monoclonal antibody hybridoma techniques for isolation of monoclonal antibodies. Alternatively, it is now possible to produce transgenic animals (e.g., mice) that are capable, upon immunization, of producing a full repertoire of human antibodies in the absence of endogenous immunoglobulin production. For example, it has been described that the homozygous deletion of the antibody heavy-chain rejoining region (JH) gene in chimeric and germ-line mutant mice results in complete inhibition of endogenous antibody production. Transfer of the human germ-line immunoglobulin gene array in such germ-line mutant mice will result in the production of human antibodies upon antigen challenge. See, e.g., Jakobovits et al., Proc. Natl. Acad. Sci. USA, 90:2551 (1993); Jakobovits et al., Nature, 362:255-258 (1993); Brugggermann et al., Year in Immunol., 73:3 (1993); and Duschosal et al., Nature 355:258 (1992).

[0064] The monoclonal antibodies herein specifically include “chimeric” antibodies (immunoglobulins) in which a portion of the heavy and/or light chain is identical with or homologous to corresponding sequences in antibodies derived from a particular species or belonging to a particular antibody class or subclass, while the remainder of the chain(s) is identical with or homologous to corresponding sequences in antibodies derived from another species or belonging to another antibody class or subclass, as well as fragments of such antibodies, so long as they exhibit the desired biological activity (U.S. Pat. No. 4,816,567; and Morrison et al., Proc. Natl. Acad. Sci. USA 81:6851-6855 (1984)).

[0065] The term “hypervariable region” when used herein refers to the amino acid residues of an antibody which are responsible for antigen-binding. The hypervariable region comprises amino acid residues from a “complementarity determining region” or “CDR” (i.e. residues 24-34 (L1), 50-56 (L2) and 89-97 (L3) in the light chain variable domain and 31-35 (H1), 50-65 (H2) and 95-102 (H3) in the heavy chain variable domain; Kabat et al., Sequences of Polypeptides of Immunological Interest, 5th Ed. Public Health Service, National Institutes of Health, Bethesda, Md. (1991)) and/or those residues from a “hypervariable loop” (i.e. residues 26-32 (L1), 50-52 (L2) and 91-95 (L3) in the light chain variable domain and 26-32 (H1), 53-55 (H2) and 96-101 (H3) in the heavy chain variable domain; Chothia and Lesk J. Mol. Biol., 196:901-917 (1987).) “Framework”
or “FR” residues are those variable domain residues other than the hypervariable region residues as herein defined.

[0066] “Humanized” forms of non-human (e.g., murine) antibodies are chimeric antibodies that contain minimal sequence derived from non-human immunoglobulin. For the most part, humanized antibodies are human immunoglobulins (recipient antibody) in which residues from a hypervariable region of the recipient are replaced by residues from a hypervariable region of a non-human species (donor antibody) such as mouse, rat, rabbit or nonhuman primate having the desired specificity, affinity, and capacity. In some instances, Fv framework region (FR) residues of the human immunoglobulin are replaced by corresponding non-human residues. Furthermore, humanized antibodies may comprise residues which are not found in the native antigen binding in the donor antibody. These modifications are made to further refine antibody performance. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the hypervariable loops correspond to those of a non-human immunoglobulin and all or substantially all of the FR regions are those of a human immunoglobulin sequence. The humanized antibody optionally also will comprise at least a portion of a human immunoglobulin constant region (Fc), typically that of a human immunoglobulin.

[0067] The choice of human variable domains, both light and heavy, to be used in making the humanized antibodies is very important to reduce antigenicity. According to the so-called “best-fit” method, the sequence of the variable domain of a rodent antibody is screened against the entire library of known human variable domain sequences. The human sequence which is closest to that of the rodent is then accepted as the human framework (FR) for the humanized antibody (Sims et al., J. Immunol., 151:2296 (1993); Chothia et al., J. Mol. Biol., 196:901 (1987)).

[0068] Another method uses a particular framework derived from the consensus sequence of all human antibodies of a particular subgroup of light or heavy chains. The same framework may be used for several different humanized antibodies (Carter et al., Proc. Natl. Acad. Sci. USA, 89:4285 (1992); Presta et al., J. Immunol., 151:2623 (1993)).

[0069] It is further important that antibodies be humanized with retention of high affinity for the antigen and other favorable biological properties. To achieve this goal, according to a preferred method, humanized antibodies are prepared by a process of analysis of the parental sequences and various humanized preparations of the models of the parental and humanized sequences. Three-dimensional immunoglobulin models are commonly available and are familiar to those skilled in the art. Computer programs are available which illustrate and display probable three-dimensional conformational structures of selected candidate immunoglobulin sequences. Inspection of these displays permits analysis of the likely role of the residues in the functioning of the candidate immunoglobulin sequence, i.e., the analysis of residues that influence the ability of the candidate immunoglobulin to bind its antigen. In this way, FR residues can be selected and combined from the recipient and import sequences so that the desired antibody characteristic, such as increased affinity for the target antigen(s), is achieved. In general, the CDR residues are directly and most substantially involved in influencing antigen binding.

[0070] “Antibody fragments” comprise a portion of a full length antibody, generally the antigen binding or variable region thereof. Examples of antibody fragments include Fab, Fab’, F(ab’)2, and Fv fragments; diabodies; linear antibodies; single-chain antibody molecules; and multispecific antibodies formed from antibody fragments. Various techniques have been developed for the production of antibody fragments. Traditionally, these fragments were derived via proteolytic digestion of intact antibodies (see, e.g., Morimoto et al., Journal of Biochemical and Biophysical Methods, 24:107-117 (1992) and Brennan et al., Science, 229:81 (1985)). However, these fragments can now be produced directly by recombinant host cells. For example, the antibody fragments can be isolated from the antibody phage libraries discussed above. Alternatively, Fab-SH fragments can be directly recovered from E. coli and chemically coupled to form F(ab’)2 fragments (Carter et al., Bio/Technology, 10:163-167 (1992)). In another embodiment, the F(ab’).2 is formed, using the leucine zipper GCN4 to promote assembly of the F(ab’)2 molecule. According to another approach, F(ab’)2 fragments can be isolated directly from recombinant host cell culture. Other techniques for the production of antibody fragments will be apparent to the skilled practitioner.

[0071] In other embodiments, the antibody of choice is a single chain Fv fragment (scFv). See WO 93/16185. “Single-chain Fv” or “scFv” antibody fragments comprise the Vµ and Vλ domains of an antibody, wherein these domains are present in a single polypeptide chain. Generally, the Fv polypeptide further comprises a polypeptide linker between the Vµ and Vλ domains which enables the scFv to form the desired structure for antigen binding. For a review of scFv see Pluckthun in The Pharmacology of Monoclonal Antibodies, vol., 113, Rosenberg and Moore eds. Springer-Verlag, New York, pp. 269-315 (1994).

[0072] The term “diabodies” refers to small antibody fragments with two antigen-binding sites, which fragments comprise a heavy chain variable domain (Vµ) connected to a light chain variable domain (Vλ) in the same polypeptide chain (Vµ-Vλ). By using a linker that is too short to allow pairing between the two domains on the same chain, the chains are forced to pair with the complementary domains of another chain and create two antigen-binding sites. Diabodies are described more fully in, for example, EP 404,087; WO 93/11161; and Hollinger et al., Proc. Natl. Acad. Sci. USA, 90:6444-6448 (1993).

[0073] The expression “linear antibodies” when used throughout this application refers to the antibodies described in Zapata et al., Polypeptide Eng., 8(10):1057-1062 (1995). Briefly, these antibodies comprise a pair of tandem Fd segments (Vµ-Cµ1-Vµ-Cµ2) which form a pair of antigen binding regions. Linear antibodies can be bispecific or monospecific.

[0074] “Multispecific antibodies” have binding specificities for at least two different epitopes, where the epitopes are usually from different antigens. While such molecules normally will only bind two antigens (i.e. bispecific antibodies, BsAbs), antibodies with additional specificities such as trispecific antibodies are encompassed by this expression when used herein. Examples of BsAbs include those with one arm directed against a tumor cell antigen and the other arm directed against a cytotoxic trigger molecule such as anti-FcγRII/anti-CD15, anti-p185/anti-FcγRIII (CD16), anti-CD3/anti-malignant B-cell (1D10), anti-CD3/anti-
p18\(^{HER2}\), anti-CD3/anti-p97, anti-CD3/anti-renal cell carcinoma, anti-CD3/anti-OVCAR-3, anti-CD3/L-D1 (anticolon carcinoma), anti-CD3/anti-melanocyte stimulating hormone analog, anti-EGF receptor/anti-CD3, anti-CD3/anti-CAM1, anti-CD3/anti-C19, anti-CD3/MoV18, anti-neural cell adhesion molecule (NCAM)/anti-CD3, anti-folate binding protein (FBP)/anti-CD3, anti-pan carcinoma associated antigen (AMOC-31)/anti-CD3, BsAbs with one arm which binds specifically to a tumor antigen and one arm which binds to a toxin such as anti-saporin/anti-Id-1, anti-CD22/anti-saporin, anti-CD7/anti-saporin, anti-CEA/anti-rin A chain, anti-interferon-\(\alpha\) (IFN-\(\alpha\))/anti-hybridoma idiotype, anti-CEA/anti-vinca alkaloid; BsAbs for converting enzyme activated prodrugs such as anti-CD30/anti-alkaline phosphatase (which catalyzes conversion of mitomycin phosphate prodrug to mitomycin alcohol); BsAbs which can be used as fibrinolytic agents such as anti-fibrin/anti-tissue plasminogen activator (tPA), anti-fibrin/anti-urokinase-type plasminogen activator (uPA); BsAbs for targeting immune complexes to cell surface receptors such as anti-low density lipoprotein (LDL)/anti-Fc receptor (e.g., FcγRI, or FcγRIII); BsAbs for use in therapy of infectious diseases such as anti-CD3/anti-herpes simplex virus (HSV), anti-Fc receptor/CD3 complex/anti-influenza, anti-FcγRII/anti-HIV; BsAbs for tumor detection in vitro or in vivo such as anti-CEA/anti-EOETUBE, anti-CEA/anti-DPTA, anti-p18\(^{HER2}\)/antigen-hapten; BsAbs as vaccine adjuncts; and BsAbs as diagnostic tools such as anti-rabbit IgG/anti-ferritin, anti-horse radish peroxidase (HRP)/anti-hormone, anti-somatostatin/anti-substance P, anti-HRP/anti-FITC, anti-CEA/anti-\(\beta\)-galactosidase. Examples of trispecific antibodies include anti-CD3/anti-CD4/anti-CD37, anti-CD3/anti-CD5/anti-CD37 and anti-CD3/anti-CD8/anti-CD37. Bispecific antibodies can be prepared as full length antibodies or antibody fragments (e.g. F(ab’)\(^2\), bispecific antibodies).

Antibodies with more than two valencies are contemplated. For example, trispecific antibodies can be prepared. Tutt et al. J. Immunol. 147: 60 (1991).

A “naked antibody” for the purposes herein is an antibody that is not conjugated to a cytotoxic moiety or radiolabel.

An “intact antibody” herein is one which comprises two antigen binding regions, and an Fc region. Preferably, the intact antibody has a functional Fc region.

“Treatment” refers to both therapeutic treatment and prophylactic or preventative measures. Those in need of treatment include those already disorder as well as those in which the disorder is to be prevented.

A “disorder” is any condition that would benefit from treatment with the antibody purified as described herein. This includes both chronic and acute disorders and diseases and those pathological conditions which predispose the mammal to the disorder in question.

The work “label” when used herein refers to a detectable compound or composition which is conjugated directly or indirectly to the antibody. The label may be itself detectable (e.g., radiotiose labels or fluorescent labels) or, in the case of an enzymatic label, may catalyze chemical alteration of a substrate compound or composition which is detectable.

The term “cytotoxic agent” as used herein refers to a substance that inhibits or prevents the function of cells and/or causes destruction of cells. The term is intended to include radioactive isotopes (e.g. Ar\(^{211}\), I\(^{131}\), I\(^{125}\), Y\(^{90}\), Re\(^{186}\), Re\(^{188}\), Sm\(^{157}\), Bi\(^{212}\), Pa\(^{223}\) and radioactive isotopes of Lu), chemotherapeutic agents, and toxins such as small-molecule toxins or enzymatically active toxins of bacterial, fungal, plant or animal origin, or fragments thereof.

Modes for Carrying Out the Invention

The invention herein provides methods for purifying an antibody from a composition (e.g. an aqueous solution) comprising the antibody and one or more contaminants. The composition is generally one resulting from the recombinant production of the antibody, but may be that resulting from production of the antibody by peptide synthesis (or other synthetic means) or the antibody may be purified from a native source of the antibody. Preferably the antibody binds human CD20 antigen, such as rituximab, or binds human VEGF antigen, such as bevacizumab.

Recombinant Production of Antibodies

For recombinant production of the antibody, the nucleic acid encoding it is isolated and inserted into a replicable vector for further cloning (amplification of the DNA) or for expression. DNA encoding the antibody is readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of the antibody). Many vectors are available. The vector components generally include, but are not limited to, one or more of the following: a signal sequence, an origin of replication, one or more marker genes, an enhancer element, a promoter, and a transcription termination sequence (e.g., as described in U.S. Pat. No. 5,534,615, specifically incorporated herein by reference).

Suitable host cells for cloning or expressing the DNA in the vectors herein are the prokaryote, yeast, or higher eukaryotic cells. Suitable prokaryotes for this purpose include eubacteria, such as Gram-negative or Gram-positive organisms, for example, Enterobacteriaceae such as Escherichia, e.g., E. coli, Enterobacter Erwinia, Klebsiella, Proteus, Salmonella, e.g., Salmonella typhimurium, Serratia, e.g., Serratia marcescens, and Shigella, as well as Bacilli such as B. subtilis and B. licheniformis (e.g., B. licheniformis 41P disclosed in DD 266,710 published 12 Apr. 1989), Pseudomonas such as P. aeruginosa, and Streptomyces. One preferred E. coli cloning host is E. coli 294 (ATCC 31,446) although other strains such as E. coli B, E. coli X1776 (ATCC 31,537), and E. coli W3110 (ATCC 27,325) are suitable. These examples are illustrative rather than limiting.

In addition to prokaryotes, eukaryotic microbes such as filamentous fungi or yeast are suitable cloning or expression hosts for antibody encoding vectors. Saccharomyces cerevisiae, or common baker’s yeast, is the most commonly used among lower eukaryotic host microorganisms. However, a number of other genera, species, and strains are commonly available and useful herein, such as Schizosaccharomyces pombe; Klyueromyces hosts such as, e.g., K. lactis, K. fragilis (ATCC 12,424), K. bulgaricus (ATCC 16,045), K. wickerhammi (ATCC 24,178), K. waltii (ATCC 56,500), K. drosophilarum (ATCC 36,906), K. thermotolerans, and K. marxianus; yarrowia (EP 402,226); Pichia pastoris (EP 183,070), Candida; Trichoderma reesia (EP 244,234); Neurospora crassa; Schwanitomyces such as
Schwanniomyces occidentalis; and filamentous fungi such as, e.g., Neurospora, Penicillium, Toxoplasma, and Aspergillus hosts such as A. nidulans and A. niger.

Suitable host cells for the expression of glycosylated antibody are derived from multicellular organisms. Examples of invertebrate cells include plant and insect cells. Numerous baculoviral strains and variants and corresponding permissive insect host cells from hosts such as Spodoptera frugiperda (caterpillar), Aedes aegypti (mosquito), Aedes albopictus (mosquito), Drosopila melanogaster (fruitfly), and Bombyx mori have been identified. A variety of viral strains for transfection are publicly available, e.g., the L-1 variant of Autographa californiae NPV and the Bm-5 strain of Bombyx mori NPV, and such viruses may be used as the virus herein according to the present invention, particularly for transfection of Spodoptera frugiperda cells. Plant cell cultures of cotton, corn, potato, soybean, petunia, tomato, and tobacco can also be utilized as hosts.

However, interest has been greatest in vertebrate cells, and propagation of vertebrate cells in culture (tissue culture have become a routine procedure. Examples of useful mammalian host cell lines include, but are not limited to, monkey kidney CV1 cells transformed by SV40 (COS-7, ATCC CRL 1651); human embryonic kidney cells (293 or 293 cells) subcloned for growth in suspension culture, Graham et al., J. Gen Virol. 36:59 (1977)); baby hamster kidney cells (BHK, ATCC CCL 10); Chinese hamster ovary cells/ DHR (CHO, Urlaub et al., Proc. Natl. Acad. Sci. USA 77:4216 (1980)); mouse sertoli cells (TM4, Mather, Biol. Reprod. 23:243-251 (1980)); monkey kidney cells (CV1 ATCC CCL 70); African green monkey kidney cells (VERO-76, ATCC CRL-1587); human cervical carcinoma cells (HELA, ATCC CCL 2); canine kidney cells (MDCK, ATCC CCL 34); buffalo rat liver cells (BRL 3A, ATCC CRL 1442); human lung cells (W138, ATCC CCL 75); human liver cells (HeP G2, HB 8065); mouse mammary tumor (MMT 060562, ATCC CCL51); TRI cells (Mather et al., Annals N.Y. Acad. Sci. 383:44-68 (1982)); MRC 5 cells; FS4 cells; and human hepatoma cells (HeP G2). Often, CHO cells are preferred for the expression of antibodies, and may be advantageously used to produce the antibodies purified in accordance with the present invention.

Host cells are transformed with the above-described expression or cloning vectors for antibody production and cultured in conventional nutrient media modified as appropriate for inducing promoters, selecting transformants, or amplifying the genes encoding the desired sequences.

The host cells used to produce the antibody of this invention may be cultured in a variety of media. Commercially available media such as Ham's F10 (Sigma), Minimal Essential Medium (MEM), RPMI-1640 (Sigma), and Dulbecco's Modified Eagle's Medium (DMEM, Sigma) are suitable for culturing the host cells. In addition, any of the media described in Ham et al., Meth. Enz. 58:44 (1979) Barnes et al., Anal. Biochem. 102:255 (1980), U.S. Pat. Nos. 4,767,704; 4,657,866; 4,927,762; 4,560,655; or 5,122,469; WO 90/03430; WO 87/00195; or U.S. Patent Re. 30,985 may be used as culture media for the host cells. Any of these media may be supplemented as necessary with hormones and/or other growth factors (such as insulin, transferrin, or epidermal growth factor), salts (such as sodium chloride, calcium, magnesium, and phosphate), buffers (such as HEPES), nucleotides (such as adenosine and thymidine), antibiotics (such as garamycin; GENTAMYCIN®), trace elements (defined as inorganic compounds usually present at final concentrations in the micromolar range), and glucose or an equivalent energy source. Any other necessary supplements may also be included at appropriate concentrations that would be known to those skilled in the art. The culture conditions, such as temperature, pH, and the like, are those previously used with the host cell selected for expression, and will be apparent to the ordinarily skilled artisan.

When using recombinant techniques, the antibody can be produced intracellularly, in the periplasmic space, or directly secreted into the medium. If the antibody is produced intracellularly, as a first step, the particulate debris, either host cells or lysed cells (e.g. resulting from homogenization), is removed, for example, by centrifugation or ultrafiltration. Where the antibody is secreted into the medium, supernatants from such expression systems may be concentrated using a commercially available protein concentration filter, for example, an Amicon or Millipore Pellicon ultrafiltration unit.

The Cation Exchange Chromatography Method of the Invention

In the preferred embodiment of the invention, the compositions to be subjected to the purification method herein is a recombinantly produced antibody, preferably an intact antibody, expressed by a Chinese Hamster Ovary (CHO) recombinant host cell culture. Optionally, the composition has been subjected to at least one purification step prior to cation exchange chromatography. The composition contains the antibody of interest and one or more contaminants, such as Chinese Hamster Ovary Proteins (CHOP); leached protein A; nucleic acid; a variant, fragment aggregate or derivative of the desired antibody; another polypeptide; endotoxin; viral contaminant; cell culture media component (e.g. garamycin; GENTAMYCIN®), etc.

Examples of additional purification procedures which may be performed prior to, during, or following the cation exchange chromatography method include fractionation on a hydrophobic interaction chromatography (e.g., on PHENYL-SEPHAROSE™), ethanol precipitation, isoelectric focusing, Reverse Phase HPLC, chromatography on silica, chromatography on HEPARIN SEPHAROSE™, anion, exchange chromatography, further cation exchange chromatography, mixed-mode ion exchange, chromatofocusing, SDS-PAGE, ammonium sulfate precipitation, hydroxyapatite chromatography, gel electrophoresis, dialysis, hydrophobic change induction, chromatography, and affinity chromatography (e.g. using protein A, protein G, an antibody, or a specific substrate, ligand or antigen as the capture reagent).

According to the present invention, the cation exchange purification scheme typically includes the following steps performed sequentially: (1) equilibration of the cation exchange material; (2) loading the composition to be purified onto the cation exchange material; (3) a first wash step; (4) a second wash step, and (5) elution of the antibody of interest.

By including at least two wash steps in the cation exchange purification scheme, at least the first of which is conducted at high pH (about pH 6.8 or greater), the efficacy of purification can be significantly improved. In particular, performing the first wash step using a wash buffer with a pH in the range from about 6.8 to about 9.0 (e.g. from about 7.0 to about 8.5)
to 8.0), such as, for example, about pH 7.8 or about pH 7.0, contaminants as described above are removed more efficiently than using the conventional lower pH range of about 5.0 to about 5.5. As a result, the host cell protein content of the composition comprising the antibody eluted from the cation exchange material is typically less than about 200 ppm, which is below the approximately 500 ppm level achieved using one wash step at a pH of about 5 to 5.5.

[0095] In the preferred embodiment of the invention, the cation exchange material comprises cross-linked poly(styrene-divinylbenzene) flow-through particles (solid phase) coated with a polyhydroxylated polymer functionalized with sulfopropyl groups, for example, a POROS 50HS® column, available from Applied Biosystems.

[0096] Usually, an equilibration buffer is passed over or through the cation exchange material prior to loading the composition comprising the antibody of interest and one or more contaminants onto the material. In the preferred embodiment of the invention, the equilibration buffer has a pH from about 5.0 to about 6.0, for example about pH 5.5. One exemplary equilibration buffer comprises 19 mM MES, 60 mM NaCl, pH 5.50. Another exemplary equilibration buffer comprises 23 mM MES, 60 mM NaCl, pH 5.50.

[0097] Following equilibration, an aqueous solution comprising the antibody of interest and one or more contaminants is loaded onto the cation exchange material. Optionally, the pH of the load is in the range from about 4.0 to about 6.0, for example about pH 5.0 or about pH 5.5. In a preferred embodiment, a conditioned product pool from a prior purification step is loaded. In one embodiment, a Protein A pool from a prior Protein A chromatography purification, pH 5.0 is loaded on the cation exchange material. In another embodiment, a conditioned Q-SEPHAROSE® pool, pH 5.5 is loaded onto the cation exchange material. Exemplary load densities are in the range from about 10 to about 100 g/L resin, preferably from about 10 to about 60 g/L resin, most preferably from about 15 to about 45 g/L resin. The antibody of interest is bound to the cation exchange material as a result of this loading step.

[0098] After loading, the cation exchange material is washed in a first wash step with a first wash buffer. During the wash process, wash buffer is passed over the cation exchange material. The composition of the wash buffer is typically chosen to elute as many contaminants as possible from the resin without eluting a substantial amount of the antibody of interest. The pH of the first wash buffer is generally higher than that of the equilibration buffer and/or the loaded composition, for example about 2 to about 3 pH units higher. Preferably the pH of the first wash buffer is in the range from about pH 6.8 to about 9.0, preferably from about pH 6.8 to about 8.0, for example about pH 7.8 or about pH 7.0. Examples of buffers which buffer in this pH range include, but are not limited to HEPES, MES, sodium acetate, TRIS/ HCl, Tris(hydroxymethyl)aminomethane hydrochloride/NaOH, Bicine/ HCl, Tricine/HCl etc. The preferred first wash buffer comprises or consists of: (1) 25 mM HEPES, pH 7.8 or (2) 25 mM MOPS, pH 7.0.

[0099] In this regard, the present invention provides a composition comprising a recombinant chimERIC CD20 antibody, such as rituximab, in 25 mM HEPES, pH 7.8. The invention also provides a recombinant humanized VEGF antibody, such as bevacizumab, in 25 mM MOPS, pH 7.0. Such compositions are useful, among other things, as intermediate compositions used in the purification of these products.

[0100] The invention herein generally entails at least one further, or a second, wash step using a second wash buffer. The pH of the second wash buffer preferably is lower than that of the first wash buffer, for example from about 2 to about 3 pH units lower. So, for example, the pH of the second wash buffer may be in the range from about pH 5.0 to about pH 6.0. Preferably, the pH of the second wash buffer is about 5.5. Examples of buffers which buffer in this pH range include, but are not limited to, MES, acetic acid/ sodium acetate or NaOH, NaH2PO4/Na2HPO4, Bis/Tris/ HCl. MES, pH 5.5 is the preferred buffer for the second wash. In one embodiment, the second wash buffer comprises or consists of: 19 mM MES, 10 mM NaCl, pH 5.50. In another embodiment, the second wash buffer comprises or consists of 23 mM MES, 10 mM NaCl, pH 5.50.

[0101] While additional wash steps may be employed, preferably only a first and second wash step are performed, prior to eluting the desired antibody. Contaminants such as those discussed above are removed from the cation exchange material during the first and/or second wash step. Preferably, the first wash step removes most of the contaminants.

[0102] Following the wash step(s) noted above, the desired antibody is eluted from the cation exchange material. Elution of the antibody may be achieved by increasing the conductivity or ionic strength. Desirably, the conductivity of the elution buffer is greater than about 10 mM. Increased conductivity may be achieved by including a relatively high salt concentration in the elution buffer. Exemplary salts for this purpose include, without limitation, sodium acetate, sodium chloride (NaCl), and potassium chloride (KCl). In one embodiment, the elution buffer comprises from about 100 to about 300 mM NaCl. The elution buffer generally will have approximately the same pH as the second wash buffer. A preferred elution buffer comprises: 10 mM MES, 160 mM NaCl, pH 5.5. Another preferred elution buffer comprises: 23 mM MES, 175 mM NaCl, pH 5.5. Elution preferably involves step elution (as opposed to gradient elution).

[0103] While the elution step is optionally followed by a regeneration step, such is not necessary according to the preferred embodiment of the invention.

[0104] While additional steps are contemplated, preferably the cation exchange purification method herein consists of only the following steps: equilibration (e.g. using equilibration buffer pH about 5.5), loading a composition comprising antibody and contaminant(s) (e.g. where pH of the loaded composition is about 5.0 or about 5.5), first wash step for eluting contaminants (e.g. using first wash buffer pH about 7.8 or first wash buffer pH about 7.0), second wash step (e.g. using second wash buffer pH about 5.5), and elution (e.g. using elution buffer pH about 5.5, and increased conductivity relative to each, of the earlier steps for eluting antibody).

[0105] The antibody preparation, obtained according to the cation exchange chromatography method herein may be subjected to additional purification steps, if necessary. Exemplary further purification steps have been discussed above.

[0106] Optionally, the antibody is conjugated to one or more heterologous molecules as desired. The heterologous
molecule may, for example, be one which increases the serum half-life of the antibody (e.g. polyethylene glycol, PEG), or it may be a label (e.g. an enzyme, fluorescent label and/or radionuclide), or a cytotoxic molecule (e.g., a toxin, chemotherapeutic drug, or radioactive isotope etc).

[0107] A therapeutic formulation comprising the antibody, optionally conjugated with a heterologous molecule, may be prepared by mixing the antibody having the desired degree of purity with optional pharmaceutically acceptable carriers, excipients or stabilizers (Remington's Pharmaceutical Sciences 16th edition, Osol, A. Ed. (1980)), in the form of lyophilized formulations or aqueous solutions. "Pharmaceutically acceptable" carriers, excipients, or stabilizers are nontoxic to the dosages and concentrations employed, and include buffers such as phosphate, citrate, and other organic acids; antioxidants including ascorbic acid and methionine; preservatives (such as octadecylmethylbenzyl ammonium chloride; hexamethonium chloride; benzalkonium chloride, benzenethonium chloride; phenol, butyl or benzyl alcohol; alkyl parabens such as methyl or propyl paraben; catechol; resorcinol; cyclohexanone; 3-pentanol; and m-cresol); low molecular weight (less than about 10 residues) polypeptide; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids such as glycine, glutamine, asparagine, histidine, arginine, or lysine; monosaccharides, disaccharides, and other carbohydrates including glucose, mannose, or dextrins; chelating agents such as EDTA; sugars such as sucrose, mannitol, trehalose or sorbitol; salt-forming counter-ions such as sodium; metal complexes (e.g., Zn-protein complexes); and/or non-ionic surfactants such as TWENT™, PLURONICS™ or polyethylene glycol (PEG).

[0108] The active ingredients may also be entrapped in microcapsule prepared, for example, by coacervation techniques or by interfacial polymerization, for example, hydroxyethylcellulose or gelatin-microcapsule and poly(methylmethacrylate) microcapsule, respectively, in colloidal drug delivery systems (for example, liposomes, albumin microspheres, microemulsions, nano-particles and nanocapsules) or in macroemulsions. Such techniques as disclosed in Remington's Pharmaceutical Sciences 16th edition, Osol, A. Ed. (1980).

[0109] The formulation to be used for in vivo administration must be sterile. This is readily accomplished by filtration through sterile filtration membranes.

[0110] Sustained-release preparations may be prepared. Suitable examples of sustained-release preparations include semipermeable matrices of solid hydrophobic polymers containing the antibody variant, which matrices are in the form of shaped articles, e.g., films, or microcapsule. Examples of sustained-release matrices include polyesters, hydrogels (for example, poly(2-hydroxyethyl-methacrylate), or poly(vinyl alcohol)), polyacrylates (U.S. Pat. No. 3,773,919), copolymers of L-glutamic acid and ethyl-L-glutamate, non-degradable ethylene-vinyl acetate, degradable lactic acid-glycolic acid copolymers such as the LUPRON DEPOT™ (injectable microspheres composed of lactic acid-glycolic acid copolymer and leuprolide acetate), and poly-D-(-)-3-hydroxybutyric acid.

[0111] The antibody purified as disclosed herein or the composition comprising the antibody and a pharmaceutically acceptable carrier is then used for various diagnostic, therapeutic or other uses known for such antibodies and compositions. For example, the antibody may be used to treat a disorder in a mammal by administering a therapeutically effective amount of the antibody to the mammal. In the case of a CD20 antibody such as rituximab it can be used to deplete B-cells, treat lymphoma (for example Non-Hodgkin's Lymphoma, NHL), or leukemia (for example Chronic Lymphocytic Leukemia, CLL) as well as autoimmune diseases such as rheumatoid arthritis (RA), multiple sclerosis (MS), lupus etc. For an antibody that binds a VEGF, such as bevacizumab, it can be used to inhibit angiogenesis, treat cancer, and treat macular degeneration, etc.

[0112] The following examples are offered by way of illustration and not by way of limitation. The disclosures of all citations in the specification are expressly incorporated herein by reference.

EXAMPLE 1

Purification of a CD20 Antibody

[0113] This example describes an improved cation exchange chromatography process for purifying a CD20 antibody, rituximab. Rituximab is used for therapy of NHL, CLL, RA, MS, etc. The structure of the Rituximab molecule is disclosed in U.S. Pat. No. 5,736,137, Anderson et al., (expressly incorporated herein by reference) as well as FIGS. 1A-1B herein. Rituximab is commercially available from Genentech, Inc.

[0114] Cation-exchange chromatography is used to further reduce the levels of CHOP, DNA, leached protein A, garamycin (GENTAMYCIN®), Rituximab aggregates, and potential viruses. Rituximab binds to the column under the load conditions. The column is then washed, eluted, regenerated/sanitized, and stored until the next use. Multiple cycles may be used to process an entire batch of affinity pool. The cation-exchange pool may be held at room temperature up to 30°C. for up to 3 days or at 5°C. for up to 7 days.

[0115] The cation-exchange resin (POROS 50 HS®, Applied Biosystems) is packed in a column to a bed height of 17-33 cm. Before the affinity pool is loaded, the cation-exchange column is purged of storage solution, with equilibration buffer. After equilibration, the affinity pool is loaded onto the column. The product binds to the column under these conditions. The column is then washed with wash 1 buffer, followed by wash 2 buffer. Rituximab is eluted from the column using a high-ionic-strength elution buffer.

[0116] A comparison of the conditions for the process of the present invention compared to the original (control) process is provided in the following table.
<table>
<thead>
<tr>
<th>TABLE 1</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Comparison of Buffers for Rituximab Cation Exchange Chromatography Processes</strong></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Phase</th>
<th>Buffer composition (original process)</th>
<th>Buffer composition (exemplified process)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre-equilibration</td>
<td>20 mM MES, 500 mM NaCl, pH 5.50</td>
<td>None</td>
</tr>
<tr>
<td>Equilibration</td>
<td>20 mM MES, 60 mM NaCl, pH 5.50</td>
<td>19 mM MES, 60 mM NaCl, pH 5.50</td>
</tr>
<tr>
<td>Load</td>
<td>Conditioned Protein A pool, pH 5.00, Load density ≤5 g/L resin</td>
<td>Conditioned Protein A pool, pH 5.00, Load density ≤5 g/L resin</td>
</tr>
<tr>
<td>Wash 1</td>
<td>2.0 mM MES, 60 mM NaCl, pH 5.50</td>
<td>25 mM HEPES, pH 7.80</td>
</tr>
<tr>
<td>Wash 2</td>
<td>None</td>
<td>19 mM MES, 10 mM NaCl, pH 5.50</td>
</tr>
<tr>
<td>Elution</td>
<td>20 mM MES, 160 mM NaCl, pH 5.50</td>
<td>19 mM MES, 160 mM NaCl, pH 5.50</td>
</tr>
<tr>
<td>Regeneration</td>
<td>20 mM MES, 500 mM NaCl, pH 5.50</td>
<td>None</td>
</tr>
<tr>
<td>Sanitization</td>
<td>0.5N NaOH</td>
<td>0.5N NaOH</td>
</tr>
<tr>
<td>Storage</td>
<td>0.1N NaOH</td>
<td>0.1N NaOH</td>
</tr>
</tbody>
</table>

The desired pH, conductivity and molarity ranges for the load and buffers in the rituximab process are provided in the following table.

<table>
<thead>
<tr>
<th>TABLE 2</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Preferred pH, Conductivity and Molarity Ranges for Rituximab Process</strong></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Buffer</th>
<th>Buffer Composition</th>
<th>Target pH</th>
<th>Preferred Buffer Molarity Range</th>
<th>Preferred Buffer pH Range</th>
<th>Allowable Conductivity Range for Buffers</th>
</tr>
</thead>
<tbody>
<tr>
<td>Equilibration</td>
<td>19 mM MES, 60 mM NaCl</td>
<td>5.5</td>
<td>14-23 mM MES</td>
<td>5.0-6.0</td>
<td>5.0-7.2 mS/cm</td>
</tr>
<tr>
<td>Load</td>
<td>Conditioned Protein A Pool</td>
<td>5.0</td>
<td>NA</td>
<td>4.5-5.5</td>
<td>2.5-5.5 mS/cm</td>
</tr>
<tr>
<td>Wash 1</td>
<td>25 mM HEPES</td>
<td>7.8</td>
<td>15-35 mM HEPES</td>
<td>7.5-8.1</td>
<td>0.5-1.5 mS/cm</td>
</tr>
<tr>
<td>Wash 2</td>
<td>19 mM MES</td>
<td>5.5</td>
<td>14-23 mM MES</td>
<td>5.0-6.0</td>
<td>0.6-2.2 mS/cm</td>
</tr>
<tr>
<td>Elution</td>
<td>10 mM NaCl</td>
<td>5.5</td>
<td>14-23 mM NaCl</td>
<td>5.3-5.7</td>
<td>13.4-17.2 mS/cm</td>
</tr>
<tr>
<td>Sanitization</td>
<td>0.5N NaOH</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Storage</td>
<td>0.1N NaOH</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
</tbody>
</table>

*Conductivity values measured with temperature compensation based on a temperature of 20°C and an alpha value of 1.71.

[0118] The exemplified process for Rituximab purification enhanced the robustness of host cell protein removal by enabling higher removal of host cell proteins in the wash phases, resulting in lower levels of host cell proteins in the product pool (elution pool) and facilitating the removal of the impurities in the subsequent downstream step. FIG. 3 illustrates the advantages of the present process in terms of host cell proteins removal.

**EXAMPLE 2**

**Purification of a VEGF Antibody**

[0119] This example describes a cation exchange chromatography process for purifying a recombinant humanized vascular endothelial growth factor antibody (rhMAb VEGF), bevacizumab. The structure of the bevacizumab molecule is disclosed in U.S. Pat. No. 7,169,901, Presta et al., expressly incorporated herein by reference. See also FIGS. 2A-2B herein. Bevacizumab is commercially available from Genetech, Inc.

[0120] This example summarizes the development studies performed on the cation exchange step for an improved bevacizumab purification process. Three cation exchange resins were evaluated in these studies: CM SEPHAROSE FAST FLOW®, SP SEPHAROSE FAST FLOW® and POROS 50HS®. The cation exchange purification processes using these three resins were evaluated with respect to: process performance (impurities removal, retrovirus removal, and step yield), product quality, process robustness and process fit at all current manufacturing sites. Based on the data generated in these studies, POROS 50HS® showed superior process performance and robustness and was selected as the cation exchange resin for the improved purification process.

[0121] Cation exchange chromatography is the final chromatography step in the purification process. It serves to remove cell culture media components (garamycin), host
cell derived impurities (CHOP, and DNA) and aggregated forms of bevacizumab. It also functions as a viral removal step.

[0122] The column is operated in a bind-and-elute mode and is performed at ambient temperature. The column uses a cation exchange resin (POROS 50HS®). The resin consists of a porous, polystyrene-divinylbenzene bead support coupled with a negatively charged functional group. The column is removed from storage by washing with equilibration buffer. The viral filtered pool will be diluted with 0.3 volumes of water for injection (WF1) to meet the conductivity limit of ≤5.5 mS/cm. The viral filtered pool is then loaded onto the equilibrated column. The product binds to the resin. After loading, the column is washed with a high pH buffer to flush the load material through the column and remove CHOP impurities. The column is then washed with a low salt buffer to lower the pH and prepare the column for elution. Product is eluted using a step elution of high salt buffer with a maximum of 7 column volumes. After elution, the column and skid are sanitized with sanitization solution (0.5 N NaOH) prior to storage in storage solution (0.1 N NaOH) until its next use.

[0123] The following table provides a description of the conditions for the bevacizumab process of the invention herein.

<table>
<thead>
<tr>
<th>Phase</th>
<th>Buffer/Solution</th>
<th>Process Parameter</th>
<th>Flow rate (cm/hr)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bed Height (cm)</td>
<td>N/A</td>
<td>30</td>
<td>N/A</td>
</tr>
<tr>
<td>Equilibration</td>
<td>23 mM MES/60 mM NaCl pH 5.5, cond. 6.9 mS/cm</td>
<td>4 CV</td>
<td>100</td>
</tr>
<tr>
<td>Load</td>
<td>Conditioned Viral Filtered (VF) Pool pH 5.5 ± 0.2, cond. ≤5.5 mS/cm</td>
<td>15-45 g bevacizumab</td>
<td>100</td>
</tr>
<tr>
<td>Wash 1</td>
<td>25 mM MOPS, pH 7.0</td>
<td>3 CV</td>
<td>100</td>
</tr>
<tr>
<td>Wash 2</td>
<td>23 mM MES/10 mM NaCl pH 5.5, cond. 1.5 mS/cm</td>
<td>3 CV</td>
<td>100</td>
</tr>
<tr>
<td>Elution</td>
<td>23 mM MES/175 mM NaCl pH 5.5, cond. 18 mS/cm</td>
<td>7 CV</td>
<td>100</td>
</tr>
<tr>
<td>Start Pooling</td>
<td>OD280 ≥ 0.5</td>
<td>OD280 ≥ 1.0</td>
<td>N/A</td>
</tr>
<tr>
<td>End Pooling</td>
<td>OD280 ≥ 0.5</td>
<td>OD280 ≥ 1.0</td>
<td>N/A</td>
</tr>
<tr>
<td>Sanitization</td>
<td>0.5N NaOH</td>
<td>3-6 CV</td>
<td>50-100</td>
</tr>
<tr>
<td>Storage</td>
<td>0.1N NaOH</td>
<td>3-6 CV</td>
<td>50-100</td>
</tr>
</tbody>
</table>

[0124] The desired pH, conductivity and molarity ranges for the load and buffers in the bevacizumab process are provided in the following table.

<table>
<thead>
<tr>
<th>Target buffer</th>
<th>Target pH range</th>
<th>Target conductivity range</th>
<th>Preferred buffer molarity range</th>
<th>Preferred buffer pH range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Equil, 23 mM MES, 60 mM NaCl</td>
<td>5.4-5.6</td>
<td>6.1-7.7 mS/cm</td>
<td>13-33 mM MES</td>
<td>5.1-5.9</td>
</tr>
<tr>
<td>Load</td>
<td>Vf pool diluted with water for injection (WF1)</td>
<td>5.3-5.7</td>
<td>≤5.5 mS/cm</td>
<td>5.2-5.8</td>
</tr>
<tr>
<td>Wash 1</td>
<td>25 mM MOPS</td>
<td>6.9-7.1</td>
<td>0.2-1.2 mS/cm</td>
<td>6.6-7.4</td>
</tr>
<tr>
<td>Wash 2</td>
<td>23 mM MES 10 mM NaCl</td>
<td>5.4-5.6</td>
<td>1.2-1.8 mS/cm</td>
<td>5.1-5.9</td>
</tr>
<tr>
<td>Elution</td>
<td>23 mM MES 175 mM NaCl</td>
<td>5.45-5.55</td>
<td>17.5-18.5 mS/cm</td>
<td>5.4-5.6</td>
</tr>
<tr>
<td>Sanitization</td>
<td>0.5N NaOH</td>
<td>NA</td>
<td>50-60 mS/cm</td>
<td>0.5N NaCl</td>
</tr>
<tr>
<td>Storage</td>
<td>0.1N NaOH</td>
<td>NA</td>
<td>17-27 mS/cm</td>
<td>0.1N NaOH</td>
</tr>
</tbody>
</table>

[0125] The present process was found to be superior to the original bevacizumab process which used a first wash buffer pH 5.5. The new process herein was able to achieve pools with lower CHOP levels, it achieved a higher step yield and was an overall more robust process to run in manufacturing.
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURES:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE:

```
Gln Val Gln Leu Gln Gln Pro Gly Ala Glu Leu Val Lys Pro Gly
1     5     10   15
Ala Ser Val Lys Met Ser Cys Lys Ala Ser Gly Tyr Thr Phe Thr
20   25     30
Ser Tyr Asn Met His Trp Val Lys Gln Thr Pro Gly Arg Gly Leu
35   40   45
Glu Trp Ile Gly Ala Ile Tyr Pro Gly Asn Gly Asp Thr Ser Tyr
50   55   60
Asn Gln Lys Phe Lys Gly Lys Ala Thr Leu Thr Ala Asp Lys Ser
65   70   75
Ser Ser Thr Ala Tyr Met Gln Leu Ser Ser Leu Thr Ser Glu Asp
80   85   90
Ser Ala Val Tyr Tyr Cys Ala Arg Ser Thr Tyr Tyr Gly Gly Asp
95  100  105
Trp Tyr Phe Asn Val Trp Gly Ala Gly Thr Thr Val Thr Val Ser
110  115  120
Ala Ala Ser Thr Lys Gly Pro Ser Val Phe Pro Leu Ala Pro Ser
125  130  135
Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala Leu Gly Cys Leu Val
140  145  150
Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser Trp Asn Ser Gly
155  160  165
Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val Leu Gln Ser
170  175  180
Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Thr Pro Ser
185  190  195
Ser Leu Gly Thr Glu Thr Tyr Ile Cys Asn Val Asn His Lys Pro
200  205  210
Ser Asn Thr Lys Val Asp Lys Ala Glu Pro Lys Ser Cys Asp
215  220  225
Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly
230  235  240
Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu
245  250  255
Met Ile Ser Arg Thr Pro Glu Val Thr Asp Gly Val Glu Val His
260  265  270
Asn Ala Lys Thr Pro Arg Glu Glu Gly Tyr Asn Ser Thr Tyr
275  280  285
Arg Val Val Ser Val Leu Thr Val Leu His Glu Asp Trp Leu Asn
290  295  300
Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala
305  310  315
Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu
320  325  330
Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp Glu Leu Thr Lys
335  340  345
```
Aan Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser
350 355 360
Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Aen
365 370 375
Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe
380 385 390
Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
395 400 405
Aam Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Ann His
410 415 420
Tyr Thr Gln Lys Ser Leu Ser Ser Leu Ser Pro Gly Lys
425 430

<210> SEQ ID NO 2
<211> LENGTH: 213
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 2
Gln Ile Val Leu Ser Gln Ser Pro Ala Ile Leu Ser Ala Ser Pro
1 5 10 15
Gly Glu Lys Val Thr Met Thr Cys Arg Ala Ser Ser Ser Val Ser
20 25 30
Tyr Ile His Trp Phe Gln Gln Lys Pro Gly Ser Ser Pro Lys Pro
35 40 45
Trp Ile Tyr Ala Thr Ser Asn Leu Ala Ser Gly Val Pro Val Arg
50 55 60
Phe Ser Gly Ser Gly Ser Gly Thr Ser Tyr Ser Leu Thr Ile Ser
65 70 75
Arg Val Glu Ala Glu Asp Ala Thr Tyr Tyr Cys Gln Gln Trp
80 85 90
Thr Ser Asn Pro Pro Thr Phe Gly Gly Thr Lys Leu Glu Ile
95 100 105
Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
110 115 120
Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu
125 130 135
Aam Aen Phe Tyr Pro Arg Glu Ala Lys Val Glu Trp Lys Val Asp
140 145 150
Aam Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
155 160 165
Aas Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu
170 175 180
Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
185 190 195
Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
200 205 210
Gly Glu Cys

<210> SEQ ID NO 3
<211> LENGTH: 121
<212> TYPE: PRT
-continued

<213> ORGANISM: Artificial sequence
<220> FEATURE:  
<223> OTHER INFORMATION: sequence is synthesized  

<400> SEQUENCE: 3

Gln Val Gln Leu Gln Gln Pro Gly Ala Glu Leu Val Lys Pro Gly  
1   5   10   15  

Ala Ser Val Lys Met Ser Cys Lys Ala Ser Gly Tyr Thr Phe Thr  
20  25   30  

Ser Tyr Asn Met His Trp Val Lys Gln Thr Pro Gly Arg Gly Leu  
35  40   45  

Glu Trp Ile Gly Ala Ile Tyr Pro Gly Asn Gly Asp Thr Ser Tyr  
50  55   60  

Asn Gln Lys Phe Lys Gly Lys Ala Thr Leu Thr Ala Asp Lys Ser  
65  70   75  

Ser Ser Thr Ala Tyr Met Gln Leu Ser Ser Leu Thr Ser Glu Asp  
80  85   90  

Ser Ala Val Tyr Tyr Cys Ala Arg Ser Thr Tyr Tyr Gly Gly Asp  
95 100 105  

Trp Tyr Phe Asn Val Trp Gly Ala Gly Thr Thr Val Thr Val Ser  
110 115 120  

Ala

<210> SEQ ID NO 4  
<211> LENGTH: 106  
<212> TYPE: PRT  
<213> ORGANISM: Artificial sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: sequence is synthesized  

<400> SEQUENCE: 4

Gln Ile Val Leu Ser Gln Ser Pro Ala Ile Leu Ser Ala Ser Pro  
1   5   10   15  

Gly Glu Lys Val Thr Met Thr Cys Arg Ala Ser Ser Ser Val Ser  
20  25   30  

Tyr Ile His Trp Phe Gln Gln Lys Pro Gly Ser Ser Pro Lys Pro  
35  40   45  

Trp Ile Tyr Ala Thr Ser Asn Leu Ala Ser Gly Val Pro Val Arg  
50  55   60  

Phe Ser Gly Ser Gly Ser Gly Thr Ser Tyr Ser Leu Thr Ile Ser  
65  70   75  

Arg Val Glu Ala Glu Asp Ala Ala Thr Tyr Tyr Cys Gln Glu Trp  
80  85   90  

Thr Ser Asn Pro Pro Thr Phe Gly Gly Gly Thr Lys Leu Glu Ile  
95 100 105  

Lys

<210> SEQ ID NO 5  
<211> LENGTH: 5  
<212> TYPE: PRT  
<213> ORGANISM: Artificial sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: sequence is synthesized  

<400> SEQUENCE: 5

Ser Tyr Asn Met His
<210> SEQ ID NO 6
<211> LENGTH: 17
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 6

Ala Ile Tyr Pro Gly Asn Gly Asp Thr Ser Tyr Asn Gln Lys Phe
1     5     10     15
Lys Gly

<210> SEQ ID NO 7
<211> LENGTH: 12
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 7

Ser Thr Tyr Tyr Gly Gly Asp Trp Tyr Phe Asn Val
5     10

<210> SEQ ID NO 8
<211> LENGTH: 10
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 8

Arg Ala Ser Ser Ser Val Ser Tyr Ile His
5     10

<210> SEQ ID NO 9
<211> LENGTH: 7
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 9

Ala Thr Ser Asn Leu Ala Ser
5

<210> SEQ ID NO 10
<211> LENGTH: 9
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 10

Gln Gln Trp Thr Ser Asn Pro Pro Thr
5

<210> SEQ ID NO 11
<211> LENGTH: 453
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized
<table>
<thead>
<tr>
<th>Sequence: 11</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glu Val Glu Leu Val Glu Ser Gly Gly Gly Leu Val Glu Pro Gly</td>
</tr>
<tr>
<td>1</td>
</tr>
<tr>
<td>Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Tyr Thr Phe Thr</td>
</tr>
<tr>
<td>20</td>
</tr>
<tr>
<td>Asn Tyr Gly Met Asn Trp Val Arg Glu Ala Pro Gly Lys Gly Leu</td>
</tr>
<tr>
<td>35</td>
</tr>
<tr>
<td>Glu Trp Val Gly Trp Ile Asn Thr Tyr Thr Gly Glu Pro Thr Tyr</td>
</tr>
<tr>
<td>50</td>
</tr>
<tr>
<td>Ala Ala Asp Phe Lys Arg Arg Phe Thr Phe Ser Leu Asp Thr Ser</td>
</tr>
<tr>
<td>65</td>
</tr>
<tr>
<td>Lys Ser Thr Ala Tyr Leu Glu Met Asn Ser Leu Arg Ala Glu Asp</td>
</tr>
<tr>
<td>80</td>
</tr>
<tr>
<td>Thr Ala Val Tyr Cys Ala Lys Tyr Pro His Tyr Tyr Gly Ser</td>
</tr>
<tr>
<td>95</td>
</tr>
<tr>
<td>Ser His Trp Tyr Phe Asp Val Trp Gly Gln Gly Thr Leu Val Thr</td>
</tr>
<tr>
<td>110</td>
</tr>
<tr>
<td>Val Ser Ser Ala Ser Thr Lys Gly Pro Ser Val Phe Pro Leu Ala</td>
</tr>
<tr>
<td>125</td>
</tr>
<tr>
<td>Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala Leu Gly Cys</td>
</tr>
<tr>
<td>140</td>
</tr>
<tr>
<td>Leu Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser Trp Asn</td>
</tr>
<tr>
<td>155</td>
</tr>
<tr>
<td>Ser Gly Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val Leu</td>
</tr>
<tr>
<td>170</td>
</tr>
<tr>
<td>Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro</td>
</tr>
<tr>
<td>185</td>
</tr>
<tr>
<td>Ser Ser Leu Gly Thr Gin Thr Tyr Ile Cys Asn Val Asn His</td>
</tr>
<tr>
<td>200</td>
</tr>
<tr>
<td>Lys Pro Ser Asn Thr Lys Val Asp Lys Lys Val Glu Pro Lys Ser</td>
</tr>
<tr>
<td>215</td>
</tr>
<tr>
<td>Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu</td>
</tr>
<tr>
<td>230</td>
</tr>
<tr>
<td>Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp</td>
</tr>
<tr>
<td>245</td>
</tr>
<tr>
<td>Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val</td>
</tr>
<tr>
<td>260</td>
</tr>
<tr>
<td>Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val</td>
</tr>
<tr>
<td>275</td>
</tr>
<tr>
<td>Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu</td>
</tr>
<tr>
<td>290</td>
</tr>
<tr>
<td>Gin Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu</td>
</tr>
<tr>
<td>305</td>
</tr>
<tr>
<td>His Gin Asp Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser</td>
</tr>
<tr>
<td>320</td>
</tr>
<tr>
<td>Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala</td>
</tr>
<tr>
<td>335</td>
</tr>
<tr>
<td>Lys Gly Gin Pro Arg Glu Pro Gin Val Tyr Thr Leu Pro Pro Ser</td>
</tr>
<tr>
<td>350</td>
</tr>
<tr>
<td>Arg Glu Glu Met Thr Lys Asn Gin Val Ser Leu Thr Cys Leu Val</td>
</tr>
</tbody>
</table>
-continued

<table>
<thead>
<tr>
<th>365</th>
<th>370</th>
<th>375</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn</td>
<td></td>
<td></td>
</tr>
<tr>
<td>380</td>
<td>385</td>
<td>390</td>
</tr>
<tr>
<td>Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp</td>
<td></td>
<td></td>
</tr>
<tr>
<td>395</td>
<td>400</td>
<td>405</td>
</tr>
<tr>
<td>Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys</td>
<td></td>
<td></td>
</tr>
<tr>
<td>410</td>
<td>415</td>
<td>420</td>
</tr>
<tr>
<td>Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His</td>
<td></td>
<td></td>
</tr>
<tr>
<td>425</td>
<td>430</td>
<td>435</td>
</tr>
<tr>
<td>Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser</td>
<td></td>
<td></td>
</tr>
<tr>
<td>440</td>
<td>445</td>
<td>450</td>
</tr>
<tr>
<td>Pro Gly Lys</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<210> SEQ ID NO 12
<211> LENGTH: 214
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 12

<table>
<thead>
<tr>
<th>1</th>
<th>5</th>
<th>10</th>
<th>15</th>
</tr>
</thead>
<tbody>
<tr>
<td>Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>25</td>
<td>30</td>
<td></td>
</tr>
<tr>
<td>Gly Asp Arg Val Thr Ile Thr Cys Ser Ala Ser Gln Asp Ile Ser</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>35</td>
<td>40</td>
<td>45</td>
<td></td>
</tr>
<tr>
<td>Asn Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>50</td>
<td>55</td>
<td>60</td>
<td></td>
</tr>
<tr>
<td>Val Leu Ile Tyr Phe Thr Ser Ser Leu His Ser Gly Val Pro Ser</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>65</td>
<td>70</td>
<td>75</td>
<td></td>
</tr>
<tr>
<td>Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>80</td>
<td>85</td>
<td>90</td>
<td></td>
</tr>
<tr>
<td>Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gin</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>95</td>
<td>100</td>
<td>105</td>
<td></td>
</tr>
<tr>
<td>Tyr Ser Thr Val Pro Trp Thr Phe Gly Gin Gly Thr Lys Val Glu</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>110</td>
<td>115</td>
<td>120</td>
<td></td>
</tr>
<tr>
<td>Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>125</td>
<td>130</td>
<td>135</td>
<td></td>
</tr>
<tr>
<td>Ser Asp Glu Gin Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>140</td>
<td>145</td>
<td>150</td>
<td></td>
</tr>
<tr>
<td>Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gin Trp Lys Val</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>155</td>
<td>160</td>
<td>165</td>
<td></td>
</tr>
<tr>
<td>Asp Asn Ala Leu Gln Ser Gly Asn Ser Gin Glu Ser Val Thr Glu</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>170</td>
<td>175</td>
<td>180</td>
<td></td>
</tr>
<tr>
<td>Glu Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>185</td>
<td>190</td>
<td>195</td>
<td></td>
</tr>
<tr>
<td>Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>200</td>
<td>205</td>
<td>210</td>
<td></td>
</tr>
<tr>
<td>Val Thr His Gin Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Arg Gly Glu Cys</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<210> SEQ ID NO 13
<211> LENGTH: 123
-continued

-Glu Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly
1  5  10  15
Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Tyr Thr Phe Thr
15 20 25  30
Asn Tyr Gly Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu
30 35 40  45
Glu Trp Val Gly Trp Ile Asn Thr Tyr Thr Gly Glu Pro Thr Tyr
45 50 55  60
Ala Ala Asp Phe Lys Arg Arg Phe Thr Phe Ser Leu Asp Thr Ser
60 65 70  75
Lys Ser Thr Ala Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp
75 80 85  90
Thr Ala Val Tyr Cys Ala Lys Tyr Pro His Tyr Tyr Gly Ser
90 95 100 105
Ser His Trp Tyr Phe Asp Val Trp Gly Gin Gly Thr Leu Val Thr
105 110 115 120
Val Ser Ser

-continued

-Glu Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
1  5  10  15
Gly Asp Arg Val Thr Ile Thr Cys Ser Ala Ser Gln Asp Ile Ser
15 20 25  30
Asn Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys
30 35 40  45
Val Leu Ile Tyr Phe Thr Ser Leu His Ser Gly Val Pro Ser
45 50 55  60
Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
60 65 70  75
Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Cys Gln Gln
75 80 85  90
Tyr Ser Thr Val Pro Trp Thr Phe Gly Gin Gly Thr Lys Val Glu
90 95 100 105
Ile Lys Arg

-continued

-Glu Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
1  5  10  15
Gly Asp Arg Val Thr Ile Thr Cys Ser Ala Ser Gln Asp Ile Ser
15 20 25  30
Asn Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys
30 35 40  45
Val Leu Ile Tyr Phe Thr Ser Leu His Ser Gly Val Pro Ser
45 50 55  60
Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
60 65 70  75
Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Cys Gln Gln
75 80 85  90
Tyr Ser Thr Val Pro Trp Thr Phe Gly Gin Gly Thr Lys Val Glu
90 95 100 105
Ile Lys Arg

-continued

-Glu Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
1  5  10  15
Gly Asp Arg Val Thr Ile Thr Cys Ser Ala Ser Gln Asp Ile Ser
15 20 25  30
Asn Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys
30 35 40  45
Val Leu Ile Tyr Phe Thr Ser Leu His Ser Gly Val Pro Ser
45 50 55  60
Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
60 65 70  75
Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Cys Gln Gln
75 80 85  90
Tyr Ser Thr Val Pro Trp Thr Phe Gly Gin Gly Thr Lys Val Glu
90 95 100 105
Ile Lys Arg

-continued

-Glu Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
1  5  10  15
Gly Asp Arg Val Thr Ile Thr Cys Ser Ala Ser Gln Asp Ile Ser
15 20 25  30
Asn Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys
30 35 40  45
Val Leu Ile Tyr Phe Thr Ser Leu His Ser Gly Val Pro Ser
45 50 55  60
Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
60 65 70  75
Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Cys Gln Gln
75 80 85  90
Tyr Ser Thr Val Pro Trp Thr Phe Gly Gin Gly Thr Lys Val Glu
90 95 100 105
Ile Lys Arg
-continued

Gly Tyr Thr Phe Thr Asn Tyr Gly Met Asn
  5
  10

<210> SEQ ID NO 16
<211> LENGTH: 17
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 16
Trp Ile Asn Thr Tyr Thr Gly Glu Pro Thr Tyr Ala Ala Asp Phe
  1
  5
  10
  15

Lys Arg

<210> SEQ ID NO 17
<211> LENGTH: 13
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 17
Pro His Tyr Tyr Gly Ser Ser His Trp Tyr Phe Asp Val
  5
  10

<210> SEQ ID NO 18
<211> LENGTH: 11
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 18
Ser Ala Ser Gln Asp Ile Ser Asn Tyr Leu Asn
  5
  10

<210> SEQ ID NO 19
<211> LENGTH: 7
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 19
Phe Thr Ser Ser Leu His Ser
  5

<210> SEQ ID NO 20
<211> LENGTH: 9
<212> TYPE: PRT
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: sequence is synthesized

<400> SEQUENCE: 20
Gln Gln Tyr Ser Thr Val Pro Trp Thr
  5
What is claimed is:
1. A method for purifying an antibody from a composition comprising the antibody and at least one contaminant, which method comprises the sequential steps of:
   (a) loading the composition onto a cation exchange material wherein the composition is at a first pH;
   (b) washing the cation exchange material with a first wash buffer at a pH which is greater than that of the composition in (a), wherein the pH of the first wash buffer is from about 6.8 to about 9.0;
   (c) washing the cation exchange material with a second wash buffer at a pH which is less than that of the first wash buffer; and
   (d) eluting the antibody from the cation exchange material with an elution buffer at a conductivity which is substantially greater than that of the second wash buffer.
2. The method of claim 1, wherein the pH of the second wash buffer and the pH of the elution buffer are approximately the same.
3. The method of claim 1 wherein the antibody binds to human CD20.
4. The method of claim 1 wherein the antibody binds to human vascular endothelial growth factor (VEGF).
5. The method of claim 1 wherein the pH of the composition in (a) is from about 4.0 to about 6.0, the pH of the first wash buffer is from about 6.8 to about 8.0, the pH of the second wash buffer is from about 5.0 to about 6.0, and the pH of the elution buffer is from about 5.0 to about 6.0.
6. The method of claim 1 wherein the conductivity of the elution buffer is from about 10 mS/cm to about 100 mS/cm.
7. The method of claim 1 wherein the elution buffer comprises about 100 to about 300 mM NaCl.
8. The method of claim 1 wherein the cation exchange material comprises cross-linked poly(styrene-divinylbenzene) flow-through particles coated with a polyhydroxylated polymer functionalized with sulfopropyl groups.
9. The method of claim 1 wherein the contaminant is selected from the group consisting of Chinese Hamster Ovary Proteins (CHO proteins), leached protein A, DNA, aggregated antibody, cell culture media component, garamycin, Chinese Hamster Ovary Protein (CHO), DNA, viral contaminant, and viral contaminant.
10. The method of claim 1 further comprising subjecting the composition comprising the antibody to one or more further purification steps either before, during, or after steps (a) through (d) so as to obtain a homogeneous preparation of the antibody.
11. The method of claim 10 further comprising conjugating the purified antibody with a heterologous molecule.
12. The method of claim 10 or 11 further comprising preparing a pharmaceutical composition by combining the homogeneous preparation of the antibody or conjugated antibody with a pharmaceutically acceptable carrier.
13. A method for purifying an antibody that binds human CD20 from a composition comprising the antibody and one or more contaminants selected from the group consisting of Chinese Hamster Ovary Proteins (CHOP), leached protein A, DNA, and aggregated CD20 antibody, which method comprises the sequential steps of:
   (a) loading the composition onto a cation exchange material wherein the composition is at a pH from about 4.0 to about 6.0;
   (b) washing the cation exchange material with a first wash buffer at a pH from about 6.8 to about 9.0;
   (c) washing the cation exchange material with a second wash buffer at a pH from about 5.0 to about 6.0; and
   (d) eluting the antibody from the cation exchange material using an elution buffer with a pH from about 5.0 to about 6.0 and a conductivity from about 10 mS/cm to about 100 mS/cm.
14. The method of claim 13 wherein the antibody is rituximab.
15. The method of claim 13 wherein the elution buffer comprises about 100 to about 300 mM NaCl.
16. A method for purifying an antibody that binds human vascular endothelial growth factor (VEGF) from a composition comprising the antibody and one or more contaminants selected from the group consisting of a cell culture media component, garamycin, Chinese Hamster Ovary Protein (CHO), DNA, viral contaminant, and aggregated VEGF antibody, which method comprises the sequential steps of:
   (a) loading the composition onto a cation exchange material wherein the composition is at a pH from about 4.0 to about 6.0;
   (b) washing the cation exchange material with a first wash buffer at a pH from about 6.8 to about 8.0;
   (c) washing the cation exchange material with a second wash buffer at a pH from about 5.0 to about 6.0; and
   (d) eluting the antibody from the cation exchange material using an elution buffer with a pH from about 5.0 to about 6.0 and a conductivity from about 10 mS/cm to about 100 mS/cm.
17. The method of claim 16 wherein the antibody is bevacizumab.
18. The method of claim 16 wherein the elution buffer comprises about 100 to about 300 mM NaCl.
19. A composition comprising rituximab in a buffer comprising about 25 mM MOPS, at a pH of about 7.8.
20. A composition comprising bevacizumab in a buffer comprising about 25 mM MOPS at a pH of about 7.0.
* * * * *