shall include this information in the license application. The manufacturer shall describe the test specifications to verify that each sublot is identical to other sublots of the lot.

(b) Final containers and dropper assemblies. (1) Final containers and dropper assemblies shall be clean.

(2) Final containers and dropper pipettes shall be colorless and sufficiently transparent to permit observation of the contents for presence of particulate matter or increased turbidity.

(c) Date of manufacture. The date of manufacture shall be the date the manufacturer begins the last entire group of potency tests.

§660.55 Labeling.

In addition to the applicable labeling requirements of §§610.62 through 610.65 and §809.10 of this chapter, and in lieu of the requirements in §§610.60 and 610.61 of this chapter, the following requirements shall be met:

(a) Potency tests for determining anti-IgG and anti-complement activity.

(b) Specificity tests, tests for heterospecific antibodies, and additional tests for nonspecific properties.

§660.52 Reference preparations.

Reference Anti-Human Globulin preparations shall be obtained from the Center for Biologics Evaluation and Research (HFM–407) (see mailing addresses in §600.2 of this chapter), and shall be used as described in the accompanying package insert for determining the potency of Anti-Human Globulin.

§660.53 Controls for serological procedures.

Red blood cells sensitized with complement shall be tested with appropriate positive and negative control antisera. All tests shall be performed in accordance with serological testing procedures approved by the Director, Center for Biologics Evaluation and Research.

§660.54 Potency tests, specificity tests, tests for heterospecific antibodies, and additional tests for nonspecific properties.

The following tests shall be performed using test procedures approved by the Director, Center for Biologics Evaluation and Research:

(a) Potency tests for determining anti-IgG and anti-complement activity.

(b) Specificity tests, tests for heterospecific antibodies, and additional tests for nonspecific properties.
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(iii) Lot number, including any sublot designations.
(iv) Expiration date.
(v) Source of the product.
(vi) Recommended storage temperature in degrees Celsius.
(vii) Volume of product.
(viii) Appropriate cautionary statement if the Anti-Human Globulin is not polyspecific. For example, "DOES NOT CONTAIN ANTIBODIES TO IMMUNOGLOBULINS" or "DOES NOT CONTAIN ANTIBODIES TO COMPLEMENT COMPONENTS."
(ix) If the final container is not enclosed in a package, all items required for a package label shall appear on the container label.

(3) Lettering size. The type size for the designation of the specific antibody on the label of a final container shall be not less than 12 point, unless otherwise approved by the Director, Center for Biologics Evaluation and Research. The prefix anti- and other parts of the name such as polyspecific may appear in smaller type.

(4) Visual inspection. When the label has been affixed to the final container, a sufficient area of the container shall remain uncovered for its full length or for no less than 5 millimeters of the lower circumference to permit inspection of the contents.

(b) Package label. The following items shall appear either on the package label or on the final container label if see-through packaging is used:
(1) Proper name of the product, and the name of the antibody or antibodies as listed in paragraph (d) of this section.
(2) Name, address (including zip code), and license number of the manufacturer.
(3) Lot number, including any sublot designations.
(4) Expiration date.
(5) Preservative(s) used and its concentration.
(6) Number of containers, if more than one.
(7) Recommended storage temperature in degrees Celsius.
(8) Source of the product.
(9) Reference to enclosed package insert.
(10) The statement: "For In Vitro Diagnostc Use."

(12) A statement of an observable indication of an alteration of the product, e.g., turbidity, color change, precipitate, that may indicate possible deterioration of the product.

(13) Appropriate cautions.

(c) Package insert. Each final container of Anti-Human Globulin shall be accompanied by a package insert meeting the requirements of §809.10 of this chapter. If two or more final containers requiring identical package inserts are placed in a single package, only one package insert per package is required.

(d) Names of antibodies.

<table>
<thead>
<tr>
<th>Antibody designation on container label</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anti-IgG, -C3d; Polyspecific.</td>
<td>Contains anti-IgG and anti-C3d (may contain other anticomplement and anti-immunoglobulin antibodies).</td>
</tr>
<tr>
<td>Anti-IgG</td>
<td>Contains anti-IgG with no anti-complement activity (not necessarily gamma chain specific).</td>
</tr>
<tr>
<td>Anti-IgG, heavy chains.</td>
<td>Contains only antibodies reactive against human gamma chains.</td>
</tr>
<tr>
<td>Anti-C3b</td>
<td>Contains only C3b antibodies with no anti-immunoglobulin activity. <strong>Note:</strong> The antibody produced in response to immunization is usually directed against the antigenic determinant which is located in the C3c subunit; some persons have called this antibody &quot;anti-C3c.&quot; In product labeling, this antibody should be designated anti-C3b.</td>
</tr>
<tr>
<td>Anti-C3d</td>
<td>Contains only C3d antibodies with no anti-immunoglobulin activity.</td>
</tr>
<tr>
<td>Anti-C4b</td>
<td>Contains only C4b antibodies with no anti-immunoglobulin activity.</td>
</tr>
<tr>
<td>Anti-C4d</td>
<td>Contains only C4d antibodies with no anti-immunoglobulin activity.</td>
</tr>
</tbody>
</table>

Anti-Human Globulin preparations may contain one or more of the antibody specificities listed in this paragraph as described in paragraph (a)(2)(i) of this section.


PART 680—ADDITIONAL STANDARDS FOR MISCELLANEOUS PRODUCTS

Sec.
680.1 Allergenic Products.
680.2 Manufacture of Allergenic Products.
680.3 Tests.