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<th>Drug</th>
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<tr>
<td>NDA 50–725</td>
<td>AUGMENTIN ‘200’ (amoxicillin; clavulanate potassium) Powder for Oral Suspension, 200 mg/5 mL; EQ 28.5 mg base/5 mL</td>
<td>Do .................................</td>
<td>May 31, 1996.</td>
</tr>
<tr>
<td></td>
<td>AUGMENTIN ‘400’ (amoxicillin; clavulanate potassium) Powder for Oral Suspension, 400 mg/5 mL; EQ 57 mg base/5 mL</td>
<td>Do .................................</td>
<td>Do.</td>
</tr>
<tr>
<td>Do</td>
<td>AUGMENTIN ‘200’ (amoxicillin; clavulanate potassium) Chewable Tablet, 200 mg; EQ 28.5 mg base</td>
<td>Do .................................</td>
<td>Do.</td>
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<tr>
<td>Do</td>
<td>AUGMENTIN ES–600 (amoxicillin; clavulanate potassium) Powder for Oral Suspension, 600 mg/5 mL; EQ 42.9 mg base/5 mL</td>
<td>SmithKline Beecham d/b/a GlaxoSmithKline, One Franklin Plaza, Philadelphia, PA 19101.</td>
<td>June 22, 2001.</td>
</tr>
</tbody>
</table>

In a letter dated November 10, 2009, GlaxoSmithKline notified FDA that the AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document, among other drug products, were being discontinued, and FDA moved the drug products to the “Discontinued Drug Product List” section of the Orange Book. Approved ANDAs for the AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document are listed in the Orange Book, and following the discontinuation of the AUGMENTIN (amoxicillin; clavulanate potassium) products, ANDAs for certain of these products were designated as the reference listed drugs to which new ANDAs should refer.

EAS Consulting Group, LLC, submitted two citizen petitions dated March 23, 2010 (FDA–2010–P–0172), and March 26, 2010 (FDA–2010–P–0177), under 21 CFR 10.30, requesting that the Agency determine whether the following products were withdrawn from sale for reasons of safety or effectiveness:

- AUGMENTIN ‘200’ (amoxicillin; clavulanate potassium) Powder for Oral Suspension, 200 mg/5 mL; EQ 28.5 mg base/5 mL;
- AUGMENTIN ‘400’ (amoxicillin; clavulanate potassium) Powder for Oral Suspension, 400 mg/5 mL; EQ 57 mg base/5 mL; and
- AUGMENTIN ES–600 (amoxicillin; clavulanate potassium) Powder for Oral Suspension, 600 mg/5 mL; EQ 42.9 mg base/5 mL.

Although the citizen petitions did not address the other AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document, those products have also been discontinued. On our own initiative, we have also determined whether those products were withdrawn for safety or effectiveness reasons.

After considering the citizen petitions and reviewing Agency records, FDA has determined under § 314.161 that the AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document were not withdrawn for reasons of safety or effectiveness. The petitioner has identified no data or other information suggesting that the AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document were withdrawn from sale for reasons of safety or effectiveness. The Agency has reviewed our files for records concerning the withdrawal of the AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document. We have also independently evaluated relevant literature and data for possible postmarketing adverse events and have found no information that would indicate that these products were withdrawn from sale for reasons of safety or effectiveness.

Accordingly, the Agency will continue to list the AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document in the “Discontinued Drug Product List” section of the Orange Book. The “Discontinued Drug Product List” delineates, among other items, drug products that have been discontinued from marketing for reasons other than safety or effectiveness. FDA will not begin procedures to withdraw approval of approved ANDAs that refer to the AUGMENTIN (amoxicillin; clavulanate potassium) products listed in this document. Additional ANDAs that refer to these products may also be approved by the Agency as long as they meet all other legal and regulatory requirements for the approval of ANDAs. If FDA determines that labeling for these drug products should be revised to meet current standards, the Agency will advise ANDA applicants to submit such labeling.

Dated: December 1, 2010.

Leslie Kux,
Acting Assistant Commissioner for Policy.
[FR Doc. 2010–30622 Filed 12–6–10; 8:45 am]
BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration
[Docket No. FDA–2010–P–0275]

Determination That GLEEVEC (Imatinib Mesylate) Capsules, 50 Milligrams and 100 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined that GLEEVEC (imatinib mesylate) Capsules, 50 milligrams (mg) and 100 mg, were not withdrawn from sale for reasons of safety or effectiveness. This determination will allow FDA to approve abbreviated new drug applications (ANDAs) for imatinib mesylate capsules, 50 mg and 100 mg, if all other legal and regulatory requirements are met.

FOR FURTHER INFORMATION CONTACT:
Rochelle Chodock Fink, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, rm. 6236, Silver Spring, MD 20993–0002, 301–796–0838.

SUPPLEMENTARY INFORMATION: In 1984, Congress enacted the Drug Price Competition and Patent Term

### Table 1—Continued

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The Food and Drug Administration (FDA) is announcing the following public workshop: Third Annual Sentinel Initiative Public Workshop. Hosted by the Engelberg Center for Health Care Reform at The Brookings Institution, this 1-day public workshop will bring together the stakeholder community for a productive discussion on a variety of topics in active medical product surveillance, including an update on Mini-Sentinel and related activities, near-term plans for FDA’s Sentinel Initiative, and opportunities for coordination with other U.S. Department of Health and Human Services efforts that use distributed systems of automated health care data.

**Date and Time:** The public workshop will be held on January 12, 2011, from 8:30 a.m. to 4:30 p.m.

**Location:** The public workshop will be held at the Renaissance Dupont Hotel, 1143 New Hampshire Ave. NW., Washington, DC 20037.

**Contact:** Kayla Garvin, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, rm. 6331, Silver Spring, MD 20993, 301–796–7578, e-mail: sentineline@fda.hhs.gov.

**Registration:** To attend the public workshop, please register at http://guest.event.com/d/hdq5r4/1Q. When registering, provide the following information: Your name, title, company or organization (if applicable), address, phone number, and e-mail address. There is no fee to register for the public workshop, but because seating is limited, registration will be on a first-come, first-served basis. A 1-hour lunch break is scheduled; however, food will not be provided. There are multiple restaurants within walking distance of the hotel where attendees can get food. If you need special accommodations due to a disability, please contact The Brookings Institution event coordinator at 202–797–4391 or e-mail: sentineline@brookings.edu at least 7 days in advance.

**Meeting Materials:** Please be advised that as soon as workshop materials are available, they will be accessible at The Brookings Institution events Web site at http://www.brookings.edu/health/events.

Dated: December 1, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–30570 Filed 12–6–10; 8:45 am]

BILLING CODE 4160–01–P