## NEWS RELEASE ALABAMA DEPARTMENT OF PUBLIC HEALTH

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## Alabama residents received contaminated NECC products

## FOR IMMEDIATE RELEASE

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The Alabama Department of Public Health has been made aware of 19 residents who have received New England Compounding Center (NECC) contaminated steroid products from procedures in Tennessee and Florida. At this time, 12 residents have been contacted and do not have symptoms of meningitis or fungal infection. One resident has died and an autopsy is pending. ADPH is actively working to contact the remaining six residents. It is likely that additional Alabamians who received treatment in other states may be identified and contacted.

At this point in FDA's investigation, the sterility of any injectable drugs produced by NECC is of significant concern. Out of an abundance of caution, patients who had procedures after May 21, 2012, that utilized an injectable NECC product should be notified by their health care practitioner about potential risk of infection. Examples of procedures include, but are not limited to, epidurals, joint injections, eye surgeries and heart surgeries.

Earlier Tuesday, CDC provided ADPH with a list of 44 Alabama health care facilities that received NECC products. ADPH is validating the list and will be contacting these facilities to ensure that NECC products have been removed from inventory and to discuss the process for contacting their patients who received the NECC products of concern.

If any patients are identified with symptoms suggestive of meningitis or other infections, they should be referred to the appropriate medical provider. In addition, a report should be made to both FDA MedWatch (1-800-332-1088 or <a href="www.fda.gov/medwatch">www.fda.gov/medwatch</a>) and to ADPH (1-800-338-7384 or <a href="www.adph.org/Extranet/Forms/Form.asp?formID=3768">www.adph.org/Extranet/Forms/Form.asp?formID=3768</a>). Once the list of Alabama facilities is validated, it will be posted at adph.org.

Patients who received a procedure at one of the identified facilities after May 21, 2012, should self-monitor for at least 3 months following the procedure. Signs and symptoms of meningitis include fever, headache, stiff neck, nausea and vomiting. Signs and symptoms of possible infections include fever, swelling, increasing pain or drainage from surgical site. Patients experiencing any of these should contact their primary care provider and the health care practitioner who performed the procedure.