PURPOSE:
To establish a procedure to approve synthetic routed guiderail blockouts for addition to the NJDOT Bureau of Material’s Qualified Products List (QPL).

REFERENCES:
New Jersey Department of Transportation Standard Specifications for Road and Bridge Construction
Section 913.01.03

PROCEDURE:
A. Manufacturer’s Request for Approval.
The Manufacturer shall request in writing for the approval of the product. Include the following information in the request:
1. The name, address, and contact information for the manufacturer.
2. The name and designation of product to be evaluated.
3. Documentation of FHWA acceptance of the blockout.
4. Documentation of NCHRP 350, test level 3 (TL-3) results.
5. Detailed drawing and specification sheet for the blockout.
6. Materials Safety Data Sheet (MSDS).

With the request, the manufacturer shall submit a sample of the synthetic routed guiderail blockout. The ME will use the samples for future reference.

MAILING ADDRESS (USPS):
Manager, Bureau of Materials (Thiokol Bldg. 4)
New Jersey Department of Transportation
P.O. Box 600
Trenton, NJ 08625-0600

B. Bureau of Materials Review.
The ME will review the information supplied by the manufacturer and will verify the product acceptance on the NCHRP 350 Report for Guardrail Offset Blocks. The product will be approved if it complies with the NCHRP 350 requirements and is FHWA approved.
PROJECT ACCEPTANCE REQUIREMENTS:
Qualification of a product and addition to the QPL does not constitute a blanket approval of the material. The Contractor for each proposed project must submit the product and source on a Materials Questionnaire as specified in Section 106. The ME will approve the product and source on a project to project basis based on the specifications for the project. The ME will sample, test and accept the material according to the applicable Section of the NJDOT Standard Specifications for Road and Bridge Construction.

DISQUALIFICATION:
The ME may remove the product from the QPL for non-conformance with specification requirements or for a documented history of poor field performance. The manufacturer shall notify the ME, in writing, of any change in product formulation. Failure to notify the ME of changes in product formulation will result in disqualification.

REQUALIFICATION:
The ME will reevaluate a product which has been disqualified and removed from the QPL only after submission of a formal request along with acceptable evidence that the problems causing the disqualification have been resolved.

The ME may require the manufacturer to requalify the product for any of the following reasons:
1. To ensure that obsolete products are not kept on the list, the ME may request written confirmation from the manufacturer that the product is still available and has not changed formulation. Failure to respond to the Bureau’s written request will result in the product being removed from the list.
2. If the formulation of the product has changed, the ME may require that the new formulation be requalified.
3. If the Department’s standard specifications change or if the referenced ASTM or AASTHO standards change, the ME may require requalification to ensure that the product meets the new specification.