Noncompliance Report Guide for Equipment
PART 573 Defect and Noncompliance Responsibility and Reports[1]

On February 8 , 2007, Maxzone Auto Parts Corp. decided that a noncompliance with Federal Motor Vehicle Safety Standard No. 108 exists in items of motor vehicle equipment listed below, and is furnishing notification to the National Highway Traffic Safety Administration in accordance with 49 CFR Part 573 Defect and Noncompliance Responsibility and Reports.

Date this report was prepared: February 16, 2007

Furnish the manufacturer's identification code for this recall (if applicable): _____

1. **Identify the full corporate name of the fabricating manufacturer/brand name/trademark owner of the recalled item of equipment. If the recalled item of equipment is imported, provide the name and mailing address of the designated agent as prescribed by 49 U.S.C. §30164.**

Maxzone Auto Parts Corp. / DEPO brand

11016 Mulberry Ave., Unit B, Fontana, CA 92337

Identify the corporate official, by name and title, whom the agency should contact with respect to this recall.

Harry Han, Product Development

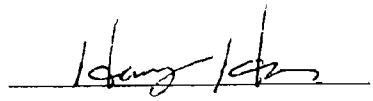
Telephone Number: : 909-822-3288 x 123 **Fax No.:** 909-822-3399

Name and Title of Person who prepared this report.

Harry Han

Product Development

Signed:



[1] Each manufacturer must furnish a report, to the Associate Administrator for Enforcement, for each noncompliance condition.

This guide was developed from 49 CFR Part 573, "Defect and Noncompliance Responsibility and Reports" and also outlines information currently requested. Any questions, please consult the complete Part 573 or contact Mr. George Person at (202) 366-5210 or by FAX at (202) 366-7882.

I. Identify the Recalled Items of Equipment

2. Identify the Items of Equipment Involved in this Recall, ***for each make and model or applicable item of equipment product line (provide illustrations or photographs as necessary to describe the item of equipment), provide*****:**

Generic name of the item:

Make: GMC **Model:** Envoy 02-06

Part Number: 335-1120L-AS **Size:**

Function: Replacement Headlamp Left Side

Other information which characterizes/distinguishes the items of equipment to be recalled:

_____

_____

Make: _____ **Model:**

Part Number: _____ **Size:**

Function:

Other information which characterizes/distinguishes the items of equipment to be recalled:

_____

_____

Make: _____ **Model:**

Part Number: _____ **Size:**

Function:

Model Years Involved:

Other information which characterizes/distinguishes the items of equipment to be recalled:

_____

_____

Make: _____ **Model:**

Part Number: _____ **Size:**

Function:

Other information which characterizes/distinguishes the items of equipment to be recalled:

_____

_____

Identify the approximate percentage of the production of all the recalled models manufactured by your company between the inclusive dates of manufacture provided above, that the recalled model population represents. For example, if the recall involved Widgets equipped with certain items of equipment from January 1, 1996, through April 1, 1997, then what was the percentage of the recalled Widgets of all Widgets manufactured during that time period.

II. Identifying the Recall Population

3. Furnish the total number of items of equipment recalled potentially containing the noncompliance.

Model	Year	Number of Items Potentially Involved
335-1120L-AS	02-06	3074pcs

Total Number Potentially Affected by the Recall: 308 pcs

4. Furnish the approximate percentage of the total number of items of equipment estimated to actually contain the noncompliance: 10.00 %

Identify and describe how the recall population was determined--in particular how the recalled models were selected and the basis for the beginning and final dates of manufacture of the recalled items of equipment:

Based on further internal testing and analysis, we've determined the subject lamps produced after Nov. 15, 2006 could potentially have been non-conforming. The last photometric testing was done on Nov. 15, 2006. The result was passed the SAE/DOT Regulation. After that, there were two lots been produced. We will issue recall notices to our distributors since we do not sell directly to end users of the subject lamps produced with the following Production Lot Numbers: 495781, 526165.

III. Describe the Noncompliance

5. Describe the noncompliance. The description should address the nature and physical location of the noncompliance. Illustrations should be provided as appropriate.

DEPO's parts failed of FMVSS108 were three position of 1.5D/2R、line test 1.5U/1R~3R and 0.5U/1R~3R.

The slight distortion of housing will change the assembly angle of reflector and influence the aiming position of photometric test. It will cause the illumination value of lamp aiming test point 1.5D/2R under the requirement. When the assembly angle of reflector deflected, that could change the aiming position and influence the values of line test 1.5U/1R~3R and 0.5U/1R~3R indirectly. The illuminate location change made above photometric test points can not pass the requirement.

Describe the cause(s) of the noncompliance condition.
Ejection system or parts release system of injection mold breakdown caused the lamp housing shape deformed. The slight distortion of housing will change the assembly angle of reflector and influence the aiming position of photometric test. It will also change the angle of incidence and illuminate location.

Describe the consequence(s) of the noncompliance condition.
Illumination value of test point 1.5D/2R、line test 1.5U/1R~3R and 0.5U/1R~3R under requirement will make the light on front of vehicle a little bit dark.

Identify any warning which can (a) precede or (b) occur.
We will do photometric test yearly to make sure our products can meet the FMVSS108 requirement.
If the test result is failed that means the lamp can't be produced.
Consumer can't identify any warning, which can precede or occur.

If the noncompliance is in a component or assembly purchased from a supplier, identify the supplier by corporate name and address.

Supplier Name : DEPO AUTO PARTS IND. CO., LTD.

Supplier Address : No. 20-3 NAN SHIH LANE LU KANG CHEN, CHANG HUA HSIEN, TAIWAN, R. O. C.

Identify the name and title of the chief executive officer or knowledgeable representative of the supplier:

Shiu-min Hsu
President

IV. Provide the Chronology in Determining the Noncompliance

6. With respect to a noncompliance, identify and provide the test results or other data (in chronological order and including dates) on which the noncompliance was determined.

095111513 test date was on Nov. 15, 2006

V. Identify the Remedy

7. A description of the manufacturer's program for remedying the noncompliance. This program shall include a plan for reimbursing an owner or purchaser who incurred costs to obtain a remedy for the problem addressed by the recall within a reasonable time in advance of the manufacturer's notification of owners, purchasers and dealers, in accordance with §573.13 of this part. A manufacturer's plan may incorporate by reference a general reimbursement plan it previously submitted to NHTSA, together with information specific to the individual recall. Information required by §573.13 that is not in a general reimbursement plan shall be submitted in the manufacturer's report to NHTSA under this section. If a manufacturer submits one or more general reimbursement plans, the manufacturer shall update each plan every two years, in accordance with §573.13. The manufacturer's remedy program and reimbursement plans will be available for inspection by the public at NHTSA headquarters.

Maxzone will issue a recall in which we will offer our customers to return the lamps for full refunds at no cost to them.

8. Furnish a description of the manufacturer's remedy for the noncompliance. Clearly describe the differences between the recall condition and the remedy.

Also we will use photometric test machine to check and make sure the remedy parts can meet FMVSS108.

We will use photometric test machine to check every recall parts. Failure parts will be scraped. There is no different between recall condition and the remedy parts.

Clearly describe the distinguishing characteristics of the remedy component/assembly versus the recalled component/assembly.

(1) We will mark "N/T" on the housing for the parts produced after correction.

(2) We will mark "R/C" on the housing for recall parts.

Identify and describe how and when the recall condition was corrected in production. If the production remedy was identical to the recall remedy in the field, so state. If the product was discontinued, so state.

By the end of February, the ejection system and parts release system of housing injection mold will be repaired. It will reduce the housing shape deformed to zero. Also we will use photometric test machine to check and make sure the production remedy parts can meet FMVSS108.

VI. Identify the Recall Schedule

9. Furnish a schedule or agenda (with specific dates) for notification to other manufacturers, dealers/retailers, and purchasers. Please, identify any foreseeable problems with implementing the recall.

Recall will be announced immediately.

VII. Furnish Recall Communications

10. Furnish a final copy of all notices, bulletins, and other communications that relate directly to the noncompliance and which are sent to more than one manufacturer, distributor, or purchaser. This includes all communications (including both original and follow-up) concerning this recall from the time your company determines the noncompliance condition on, not just the initial notification. ***A DRAFT copy of the notification documents should be submitted to the Office of Defects Investigation by Fax (202-366-7882) for review prior to mailing.***

Note: These documents are to be submitted separately from those provided in accordance with Part 579.5 requirements.

A copy of the completed Part 573 report guide should be faxed to:

Mr. Samuel Campbell at (202) 366-7097