

Safety Defect and Noncompliance Report Guide for Vehicles PART 573 Defect and Noncompliance Responsibility and Reports¹

On April 01, 2010, Main Mobility, Inc. [MFR] decided that (a defect which relates to motor vehicle safety) exists in the motor vehicles 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: April 13, 2012

Furnish the manufacturer's identification code for this recall (if applicable): 12E002

1. Identify the full corporate name of the fabricating manufacturer of the vehicle being recalled. If the recalled vehicle is imported, provide the name and mailing address of the designated agent as prescribed by 49 U.S.C. §30164.

Main Mobility, Inc.

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

Thomas McGraw, General Manager

Telephone Number: (716) 759-6811 **Fax No.:** (716) 759-6812

Name and Title of Person who prepared this report.

Thomas McGraw

General Manager

Signed:

¹ Each manufacturer must furnish a report, to the Associate Administrator for Enforcement, for each defect or noncompliance condition which relates to motor vehicle safety.

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, by FAX at (202) 366-7882, or by E-Mail to RMD.ODI@dot.gov.



I. Identify the Vehicle Models Involved in the Recall

2. Identify the Vehicles Involved in the Recall, for each make and model or applicable vehicle line (provide illustrations or photographs as necessary to describe the vehicle), provide:

Make(s): Chevrolet Model Years Involved: 2008-2010 Model(s): Express 3500			
Production Dates: Beginning: November 20, 2006 Ending: Present			
VIN Range: Beginning: <u>1GAHG39K491</u> Ending: <u>1GBHG39KX91</u>			
Vehicle Type: <u>VAN</u> Bodystyle: <u>VAN</u>			
Descriptive information which characterizes/distinguishes the recalled vehicles from those model vehicles			
not included in the recall:			
Has Braun wheelchair lift installed			
Make(s): <u>GMC</u> Model Years Involved: <u>2009</u> Model(s): <u>SAVANA 3500</u>			
Production Dates: Beginning: Ending:			
VIN Range: Beginning: <u>1GDGG25K79</u> _Ending: <u>1GDGG25K79</u>			
Vehicle Type: <u>VAN</u> Bodystyle: <u>VAN</u>			
Descriptive information which characterizes/distinguishes the recalled vehicles from those model vehicles			
not included in the recall:			
Has Braun wheelchair lift installed			
Make(s): FORD Model Years Involved: 2010 Model(s): TRANSIT CONNECT			
Production Dates: Beginning: Ending:			
VIN Range: Beginning: <u>NM0KS9AN3AT</u> Ending: <u>NM0KS9AN3AT</u>			
Vehicle Type: <u>VAN</u> Bodystyle: <u>VAN</u>			

Descriptive information which characterizes/distinguishes the recalled vehicles from those model vehicles not included in the recall:

Has Braun wheelchair lift installed



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 Vehicles equipped with certain items of equipment from January 1, 1996 through April 1, 1997, then what was the percentage of the recalled Vehicles of all Vehicles manufactured during that time period. <u>6%</u>

II. Identify the Recall Population

3. Furnish the total number of vehicles recalled potentially containing the defect or noncompliance.

Model	Year	Number of Vehicles Potentially Involved
Chevrolet Express	2008	16
Chevrolet Express	2009	9
Chevrolet Express	2010	7
GMC SAVANAss	2009	1
FORD TRANSIT CONNECT	2010	1

Total Number Potentially Affected by the Recall:

34

4. Furnish the approximate percentage of the total number of vehicles estimated to actually contain the defect or noncompliance: <u>100%</u>

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 vehicles:

The recall was determined based upon the recall of the Braun Corporation, manufacturer of the wheelchair lifts installed in these vehicles. The recalled models were selected as they are the only models on which Main Mobility builds wheelchair conversions. The dates of manufacture were determined from the start date of production of the recalled wheelchair lists to present.



III. Describe the Defect or Noncompliance

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

See Braun Recall 12E002.

Describe the cause(s) of the defect or noncompliance condition.

See Braun Recall 12E002.

Describe the consequence(s) of the defect or noncompliance condition.

See Braun Recall 12E002.

Identify any warning which can (a) precede or (b) occur.

See Braun Recall 12E002.

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

See Braun Recall 12E002.

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

See Braun Recall 12E002.

IV. Provide the Chronology in Determining the Defect/Noncompliance

If the recall is for a defect, complete item 6, otherwise item 7.

6. With respect to a defect, furnish a chronological summary (including dates) of all the principle events that were the basis for the determination of the defect. The summary should include, but not be limited to, the number of reports, accidents, injuries, fatalities, and warranty claims.

7. 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.



See Braun Recall 12E002.

V. Identify the Remedy

8. A description of the manufacturer's program for remedying the defect or 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.

See Braun Recall 12E002.

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

See Braun Recall 12E002.

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

See Braun Recall 12E002.

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.

See Braun Recall 12E002.

VI. Identify the Recall Schedule

10. 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.



See Braun Recall 12E002.

VII. Furnish Recall Communications

11. Furnish a final copy of all notices, bulletins, and other communications that relate directly to the defect or 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 defect or noncompliance condition on, not just the initial notification. A DRAFT copy of the notification documents should be submitted to this office by Fax (202-366-7882) or by E-Mail to <u>RMD.ODI@dot.gov</u> for review prior to mailing.

<u>Note</u> that these documents are to be submitted separately from those provided in accordance with Part 579.5 requirements.