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By Recall Management Division at 3:09 pm, Jun 22, 2012

MOLLY CORPORATION
Manufacture & Sales of MOLLY TROLLEY
60 Willie Hill Road P.O. Box 1799
WELLS, MAINE 04090
(207) 646-5908 FAX (207) 646-6497

12V-292
(6 Pages)

Mr, Alex Ansley
US DOT – NHTSA
Office of Defects Investigations
Recall Management / W46-412
1200 New Jersey Avenue SE
Washington, DC 20590

June 22, 2012

RE: PART 573 Defect and Noncompliance Responsibility and Report

On June 22, 2012 Molly Corporation was notified by the FTA pursuant to 49 C.F.R. Part 573, Braun Corporation has reported a safety defect in certain Century-2 and Vista-2 wheelchair lifts that were used in Molly Corporation's "Molly Trolleys". The above safety defect has been issued Safety Recall EQ12-005.

Molly Corporation Contact Information:

Jamie Bradish, Warranty Department
Molly Corporation
60 Willie Hill Road
PO Box 1799
Wells, ME 04090

Tel. (207) 646-5908
Fax. (207) 646-6497

Signed:

I. Identify the Vehicle Models Involved in the Recall

Owner	VIN #
Boston Trolley Tours, LLC **100 Terminal Street** **Charlestown, MA 02129**	**4UZAB0DTXAC.** **4UZAB0DT1AC.**

II. Identify the Recall Population

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

Vehicles Listed Above

Total Number Potentially Affected by the Recall: **2 Molly Trolleys**

4. Furnish the approximate percentage of the total number of vehicles estimated to actually contain the defect or noncompliance:

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:

See Braun Safety Recall Notice 12E-002

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 Safety Recall Notice 12E-002

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

See Braun Safety Recall Notice 12E-002

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

See Braun Safety Recall Notice 12E-002

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

See Braun Safety Recall Notice 12E-002

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 Safety Recall Notice 12E-002

Identify the name and title of the chief executive officer or knowledgeable representative of the supplier:
See Braun Safety Recall Notice 12E-002

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 Safety Recall Notice 12E-002

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 Safety Recall Notice 12E-002

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 Safety Recall Notice 12E-002

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

See Braun Safety Recall Notice 12E-002

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 Safety Recall Notice 12E-002

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 Safety Recall Notice 12E-002

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 RMD.ODI@dot.gov for review prior to mailing.***

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

Sincerely,

6/22/12

Jamie Bradish
Molly Corporation