## URGENT FIELD SAFETY NOTICE UPDATE



## Date of Letter Deployment

GE HealthCare Ref. # 34134-ROW

To: Director of Respiratory Health Care Administrator / Risk Manager Director of Biomedical / Clinical Engineering

### RE: Update to GE HealthCare's communication of potential elevated levels of formaldehyde from EVair and EVair 03 (Jun-Air) Compressors when used with the CARESCAPE R860 or Engström Carestation/Pro ventilators

Safety Based on preliminary testing, GE HealthCare previously distributed an Urgent Medical Device Correction letter to inform you of the potential for the presence of elevated levels of formaldehyde when the EVair or EVair 03 (Jun-Air) optional compressors are used with the CARESCAPE R860 or Engström Carestation/Pro ventilators, respectively. A copy of the previous letter (Ref. # 34134) is attached for your reference (Attachment 1). As communicated in the previous letter, the test conditions were not representative of typical use conditions.

We have now completed comprehensive testing and are providing an update to the previous letter in this communication.

# UPDATE: Final testing has demonstrated that there are no hazardous levels of formaldehyde when using the EVair Compressor, even at worst-case conditions of

- 40°C (104°F),
- the lowest flow condition of 2 L/min, and
- minimum bias flow.

Our comprehensive investigation of the preliminary test conditions that resulted in the issuance of the earlier precautionary letter concluded that those preliminary tests were conducted in incorrect test conditions, which led to inaccurate formaldehyde results.

While the same improper test conditions were used for the preliminary testing of the EVair 03 (Jun-Air) Compressors, they have not been manufactured for more than seven years and there are no new/unused devices available for final testing.

GE HealthCare has not received any reports of patient injury or adverse effects related to potential exposure to formaldehyde from the use of the compressors with ventilators.

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## EVair 03 (Jun-Air) Compressors:

GE HealthCare has previously communicated End of Service Support for these compressors. If you choose to continue to use these, the instructions provided in the previous letter continue to apply. As stated above, while improper test conditions were also used for the testing that led to the issuance of restrictions for use of these Compressors, since these compressors have not been manufactured for several years, GE HealthCare is unable to conduct testing at appropriate conditions. As such, the restrictions previously communicated in the attached letter (also provided below) still apply to EVair 03 (Jun-Air):

- 1. GE HealthCare recommends that the EVair 03 (Jun-Air) compressors are not used to supply air to ventilators for neonatal and infant patients (0-2 years of age).
- 2. GE HealthCare recommends that these compressors are used at a maximum room air temperature of 30°C (86°F).

Please ensure all potential users in your facility are made aware of this updated safety notification and the recommended actions.

 
 Affected
 EVair Compressor (M1230849; M1230847; GTIN: 76402146418924, 00195278366078, 07640214641892, 07640149381030, 00195278366061, 07640214641854, 07640149381023, 76402146418542) used with CARESCAPE R860 ventilator

EVair 03 (Jun-Air) Compressor (1609000; 1609002; GTIN: Not Applicable) used with Engstrom Carestation/Pro ventilators

Intended Use for EVair:

The EVair medical air compressor (EVair) is intended to be connected to a Datex- Ohmeda Inc. critical care ventilator [CARESCAPE R860] as a supply of compressed medical breathing air (compressed air). The ventilator must be operated with at least one additional supply of compressed medical breathing air or oxygen besides the EVair.

Intended Use for EVair 03 (Jun-Air):

The EVair 03 compressor is intended for use as an optional accessory to Datex- Ohmeda critical care ventilators [Engstrom Carestation/Pro] as a breathable compressed air supply. If the compressor is the primary air supply to the system, ensure that a compressed oxygen supply is also connected.

**Product** Please replace the previously provided addendum with the new attached **Correction** addendum (attachment 2).

**Contact** If you have any questions or concerns regarding this notification, please **Information** contact GE HealthCare Service or your local Service Representative.

GE Healthcare confirms that this notice has been notified to the appropriate Regulatory Agency.

Please be assured that maintaining a high level of safety and quality is our highest priority. If you have any questions, please contact us per the contact information above.

Sincerely,

Laila Gurney Chief Quality & Regulatory Officer GE HealthCare

Scott Kelley Chief Medical & Safety Officer GE HealthCare