

Drägerwerk AG & Co. KGaA, 23542 Lübeck

**To the customers and users of the  
emergency breathing devices  
Oxylog 3000, Oxylog 3000 plus and  
Oxylog 2000 plus**

November 2015

**Important safety information!!!**

**Oxylog 3000, Oxylog 3000 plus and Oxylog 2000 plus  
Failure of ventilation function with "Poti unplugged" error message**

Dear Madam / Sir,

As part of our product monitoring we have become aware of situations where the error message "Poti unplugged" was generated. In these cases, an acoustic and visual alarm is generated, the breathing system release pressure and the ventilation function stops operating. Personal injury was not reported in any of these situations.

Our investigations showed that the error message is caused by increased electrical contact resistance of the controllers (control knobs). The increased resistance is the result of an oxide layer on the controllers, which accumulates over a longer period of time. This oxide layer can only accumulate if the controllers are moved rarely or never. Our product monitoring has shown that some users rarely use the FiO2 controller or do not use it at all.

**We therefore highly recommend as a preventive action to clean any oxide layer, once. For doing this, please switch off the device and move all controllers at least 10 times from the left stop to the right stop (minimum and maximum values). Thereafter, as an ongoing measure, during every pre-use check, turn all controllers once from the left stop to the right stop.**

In particular with the error "Device malfunction – Poti unplugged", this method can also be used to put the device back into operation.

This letter includes a **supplement to your instructions for use.**

Drägerwerk AG & Co. KGaA  
Moisinger Allee 53-55  
23558 Lübeck, Germany  
Postal address:  
23542 Lübeck, Germany  
Tel. +49 451 882-0  
Fax +49 451 882-2080  
info@draeger.com  
www.draeger.com  
VAT no. DE135082211

Bank information:  
Commerzbank AG, Lübeck  
IBAN: DE95 2304 0022 0014 6795 00  
Swift-Code: COBA DE FF 230  
Sparkasse zu Lübeck  
IBAN: DE15 2305 0101 0001 0711 17  
Swift-Code: NOLADE21SPL

Company headquarters: Lübeck  
Commercial register:  
Municipal Court Lübeck HRB 7903 HL  
General partner: Drägerwerk Verwaltungs  
AG  
Company headquarters: Lübeck  
Commercial register:  
Municipal Court Lübeck HRB 7395 HL

Chairman of the Executive Board  
of  
Drägerwerk AG & Co. KGaA and  
Drägerwerk Verwaltungs AG:  
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Anton Schrofner

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This supplement expands the test for operational readiness in such a way that all controllers must be operated once every time the device is put into operation.

May we remind you that the Oxylog must only be used after verifying its operational readiness. Please include this supplement with your instructions for use and inform all affected users in your hospital.

We regret any inconvenience this information may cause but consider it necessary as a preventive measure to increase patient and user safety.

We thank you for your support.

With best regards,



Rainie Papadopoulos  
Country Quality Manager, Sales & Service  
Dräger UK & Ireland

Annex:

- Supplement instructions for use