

Ministry of Health, Welfare and Sport  
Public Health Supervisory Service  
Health Care Inspectorate  
P.O. Box 16119  
2500 BC The Hague  
The Netherlands

## STATEMENT

The undersigned, J. Moleveld, Inspector for Medical Technology in The Netherlands, herewith declares that according to the Decree on Medical Devices, which is based on the European Directive 93/42/EEC concerning medical devices,

Manufacturer:

**Fixus B.V.**  
**Frankeneng 24**  
**6716 AA Ede**  
**THE NETHERLANDS**

is authorized to manufacture and supply the medical devices mentioned below:

- 22 - Standard External Fixator for broken bones in fingers**
- 22-V External Fixator for broken bones in fingers  
with Three Dimensional Dynamic System**
- 33 - Standard External Fixator for broken bones with V-configuration**
- 33-W External Fixator for broken wrist  
with Three Dimensional Dynamic System**
- 66 - Standard External Fixator for *inter alia* Tibia**
- 66-C External Fixator with Distraction/Compression System**
- 99 - Standard External Fixator for heavy patients**
- 99-C External Fixator with Distraction/Compression System  
for inter alia leg lengthening**
- 99-T External Fixator with components for a three-point configuration**

These devices are sold freely on the Dutch market.

The present statement is drawn up at the request of the interested party in order to be submitted to the Health Authorities of EGYPT.

This statement is valid until August 17, 2009.

The Hague, August 25, 2006.

The Inspector for Medical Technology  
in The Netherlands,

J. Moleveld



Our reference : FMT/MT-U 06-30915

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