ORDOS Technologies Inc.
EMC & Safety Test Labs

Test Certificate

#5ALF880CX

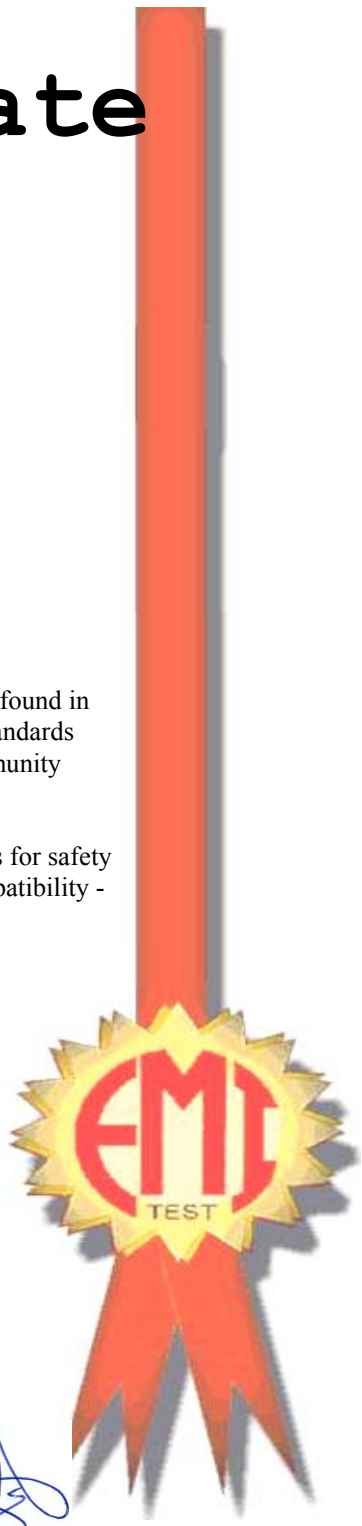
Date:	**May 2005**
Manufacturer's Name:	**Alfa Tech Medical Systems Ltd.**
Manufacturer's Address:	**Kaf -Tet be November 15/2 Bat-Yam Israel**
Equipment Under Test:	**DU 857**
Compliance with the European directives	**COUNCIL DIRECTIVE 93/42/EEC**

According to testing performed at ORDOS/E.M.I TEST LABs., the **DU 857** System was found in compliance with the Safety and EMC requirements defined in the International generic standards IEC60601-1:1988 and IEC60601-1-2:2001 which identify the Safety ,Emmision and Immunity standard requirements for Medical equipment as follows:

IEC60601-1 : 1988 +Amd(1995)	Medical electrical equipment. General requirements for safety
IEC60601-1-2 : 2001	Medical electrical equipment.Electromagnetic compatibility - Requirements and tests
IEC60601-2-5 : 2000 (Ed2)	Particular requirements for the safety of ultrasonic physiotherapy equipment
IEC60601-2-10: 1987+Amd(2002)	Particular requirements for the safety of nerve and muscle stimulators

Results and test conditions of EMC and Safety tests are specified in "ORDOS" test reports No. 5ALF880SI and 4ALF721MC. We, the undersigned, hereby declare that the equipment tested at "ORDOS/E.M.I TESTS LABS" as specified above conforms to the requirements of above European Standards.

It is the manufacturer's responsibility to assure that additional production units of this model are manufactured with identical electrical and mechanical characteristics.



Mr. Ilan Feferberg
Alfa Tech Medical Systems Ltd.

Yossi Ben-David M.Sc.
ORDOS & EMI Test Lab Manager