



**MINISTRY OF HEALTH
ECONOMIC BRANCH**

DEPARTMENT FOR MEDICAL DEVICES

1-3 Cristian Popișteanu Street, Sector 1, 010024, BUCHAREST
Telephone: 021 307 25 84; Fax: 021 307 25 85

**REGISTRATION CERTIFICATE FOR MEDICAL DEVICES
no. RO/IVD-113-295 of 24.11.2010**

Pursuant to the provisions of Government Decision no. 144/2010 on the organisation and operation of the Ministry of Health, of the minister of health's Order no. 253/2010 on registration of medical devices, of art. 29 of Government Decision no. 798/2003 on establishing the conditions for marketing and use of in vitro diagnostic medical devices and based on the documentation submitted, the Ministry of Health entered in the database the following medical devices:

NAME OF DEVICE	CODE	REGISTRATION CODE	CLASS
SEE NOW AMPHETAMINE RAPID TEST	SN 7.1	RO/IVD-113-295-01	IN VITRO OTHER
SEE NOW BARBITURATE RAPID TEST	SN 7.2	RO/IVD-113-295-02	IN VITRO OTHER
SEE NOW BENZODIAZEPINE RAPID TEST	SN 7.3	RO/IVD-113-295-03	IN VITRO OTHER
SEE NOW COCAINE RAPID TEST	SN 7.4	RO/IVD-113-295-04	IN VITRO OTHER
SEE NOW METHYLENEDIOXYMETH-AMPHETAMINE RAPID TEST	SN 7.5	RO/IVD-113-295-05	IN VITRO OTHER
SEE NOW METHAMPHETAMINE RAPID TEST	SN 7.6	RO/IVD-113-295-06	IN VITRO OTHER
SEE NOW MORPHINE RAPID TEST	SN 7.7	RO/IVD-113-295-07	IN VITRO OTHER
SEE NOW METHADONE RAPID TEST	SN 7.8	RO/IVD-113-295-08	IN VITRO OTHER
SEE NOW OPIATE RAPID TEST	SN 7.9	RO/IVD-113-295-09	IN VITRO OTHER
SEE NOW TRICYCLIC ANTIDEPRESSANTS RAPID TEST	SN 7.10	RO/IVD-113-295-10	IN VITRO OTHER
SEE NOW MARIJUANA RAPID TEST	SN 7.11	RO/IVD-113-295-11	IN VITRO OTHER
SEE NOW PHENCICLIDINE RAPID TEST	SN 7.12	RO/IVD-113-295-12	IN VITRO OTHER

put on the market according to the provisions of Government Decision no. 798/2003 by the producer:

SC CAMP MEDICA SRL

headquartered in Bucharest, 80 Vultureni Street, sector 4.

This registration was made based on the producer's declarations and it does not represent an approval or authorisation of the competent authority.

By applying the CE marking, the producer takes responsibility for compliance of the product with all applicable requirements provided in Government Decision no. 798/2003



The producer has the obligation to inform the Ministry of Health of any changes in the registered data, including suspension of the marketing of devices.

MANAGING DIRECTOR,

EC. Cristina BARBUTA

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Stamp: Romania, Ministry of Health, Economic Branch

