

REGISTRATION CERTIFICATE FOR MEDICAL DEVICES
no. RO/IVD-113-215 of 5.03.2009

Pursuant to the provisions of Government Decision no. 1718/2008 on the organisation and operation of the Ministry of Health, of the minister of public health's Order no. 92/2007 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:

"SEE NOW" RAPID TESTS

Type according to the annex.

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.

DIRECTOR,

Eng. Rodica Nadolu

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**ANNEX TO REGISTRATION CERTIFICATE no. RO/IVD-113-215
of 5.03.2009**

Official stamp Romania, Ministry of Health

"SEE NOW" RAPID TESTS

NAME OF DEVICE	TYPE	CODE	REGISTRATION CODE
See Now hCG (human Chorionic Gonadotropin) rapid test	urine/serum - strip	SN 1.1	RO/IVD-113-215-01
See Now hCG (human Chorionic Gonadotropin) rapid test	urine/serum - cassette	SN 1.2	RO/IVD-113-215-02
See Now LH (Luteinising Hormone) rapid test	urine - strip	SN 1.3	RO/IVD-113-215-03
See Now LH (Luteinising Hormone) rapid test	urine - cassette	SN 1.4	RO/IVD-113-215-04
See Now FSH (Follicle Stimulating Hormone) rapid test	urine - strip	SN 1.5	RO/IVD-113-215-05
See Now FSH (Follicle Stimulating Hormone) rapid test	urine - cassette	SN 1.6	RO/IVD-113-215-06
See Now Helicobacter Pylori rapid test	serum/plasma - strip	SN 2.1	RO/IVD-113-215-07
See Now Helicobacter Pylori rapid test	serum/plasma - cassette	SN 2.2	RO/IVD-113-215-08
See Now AFP (Alpha Fetoprotein) rapid test	serum/plasma - strip	SN 3.1	RO/IVD-113-215-09
See Now AFP (Alpha Fetoprotein) rapid test	serum/plasma - cassette	SN 3.2	RO/IVD-113-215-10
See Now CEA (Carcinoma Embryonic Antigen) rapid test	serum/plasma - strip	SN 3.3	RO/IVD-113-215-11
See Now CEA (Carcinoma Embryonic Antigen) rapid test	serum/plasma - cassette	SN 3.4	RO/IVD-113-215-12
See Now Syphilis rapid test	serum/plasma - strip	SN 4.1	RO/IVD-113-215-13
See Now Syphilis rapid test	serum/plasma - cassette	SN 4.2	RO/IVD-113-215-14
See Now Gonorrhoea rapid test	urine/urogenital secretory specimen - strip	SN 4.3	RO/IVD-113-215-15
See Now Gonorrhoea rapid test	urine/urogenital secretory specimen - cassette	SN 4.4	RO/IVD-113-215-16



**ANNEX NO. 1 of 16.07.2009 to the REGISTRATION CERTIFICATE FOR
MEDICAL DEVICES no. RO/IVD-113-215 of 5.03.2009**

Pursuant to the provisions of Government Decision no. 1718/2008 on the organisation and operation of the Ministry of Health, of the minister of public health's Order no. 92/2007 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:

The certificate is supplemented with the following medical devices:

NAME OF DEVICE	TYPE	CODE	REGISTRATION CODE
See Now Troponin I rapid test	serum/ plasma/ whole blood - cassette	SN 6.1	RO/IVD-113-215-17
See Now Myoglobin rapid test	serum/ plasma/ whole blood - cassette	SN 6.2	RO/IVD-113-215-18

The abovementioned medical devices are 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.

DIRECTOR,

Eng. Alexandru Steriu

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