VERŻJONI ELETTRONIKA

Suppliment tal-Gazzetta tal-Gvern ta' Malta, Nru. 18,983, 30 ta' Ottubru, 2012
Taqsima B

L.N. 371 of 2012

PRODUCT SAFETY ACT (CAP. 427)

In Vitro Diagnostic Medical Devices (Amendment) Regulations, 2012

IN exercise of the powers conferred by articles 38 and 39 of the Product Safety Act, the Minister for Fair Competition, Small Business and Consumers, on the advice of the Malta Competition and Consumer Affairs Authority, has made the following regulations:-

Citation, scope and commencement. S.L. 427.16

1. (1) The title of these regulations is the In Vitro Diagnostic Medical Devices (Amendment) Regulations, 2012 and these regulations shall be read and construed as one with the In Vitro Diagnostic Medical Devices, hereinafter referred to as "the principal regulations".

(2) These regulations aim to transpose Commission Directive 2011/100/EU of 20th December 2011 amending Directive 98/79/EC of the European Parliament and of the Council on *in vitro* diagnostic medical devices.

(3) These regulations shall enter into force on 1st July, 2012.

Amends ScheduleII to the principal regulations.

2. In List A of Schedule II to the principal regulations, immediately after the words "HTLV I and II, and hepatitis B, C and D", there shall be added the following:

" - Variant Creutzfeldt-Jakob disease (vCJD) assays for blood screening, diagnosis and confirmation".

Ippubblikat mid-Dipartiment tal-Informazzjoni (doi.gov.mt) — Valletta — ***Published by the Department of Information (doi.gov.mt) — Valletta***
Mitbugħ fl-Istamperija tal-Gvern fuq karta riċiklata — ***Printed at the Government Printing Press on recycled paper***

Prezz/*Price*
€0.09