Announcements 05/11/MDS-AN009

SUBJECT: Update of the Implementing Rule on LICENSING OF AUTHORISED REPRESENTATIVES (MDS-IR5) and the Guidance for Medical Device Authorised Representatives (MDS – G3).

ADDRESSES: Authorised Representatives of medical devices for both local and overseas manufacturers.

New updated version of the Implementing Rule on LICENSING OF AUTHORISED REPRESENTATIVES (MDS-IR5) and Guidance for Medical Device Authorised Representatives G3 are now published in the SFDA website under Regulations & Guidelines Tap.

Version Date :

Document Name	Version number	Version date
LICENSING OF AUTHORISED REPRESENTATIVES (MDS-IR5)	2	5th June 2011
Guidance for Medical Device Authorised Representatives (MDS – G3)	2	5th June 2011