EUDAMED Mandate Summary document that a non-EU manufacturer should provide in its Actor registration request

This document is only for the EUDAMED registration

Mandate Summary template for the registration in EUDAMED

Manufacturer name	*MF name*
Manufacturer address	*MF address*
Authorised Representative SRN	*AR SRN*
Authorised Representative Name	*AR name*
Authorised Representative Address	*AR address*
Start date of mandate	*Start date*
End date of mandate	*End date if end date is defined*
Mandated for vigilance	*[yes/no]*
Generic device group(s) source definition	*Indicate the generic device group(s) source definition e.g.:* *EMDN code(s)* *GMDN code(s)* *MF list* *AR list* *….*
List of generic device group(s) covered by this mandate	
…..	