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	*Name of the non-EU manufacturer*
Manufacturer address	*Address of the non-EU manufacturer*
Authorised Representative Actor ID/SRN	*Authorised Representative Actor ID or single registration number*
Authorised Representative Name	*Name of the Authorised Representative*
Authorised Representative Address	*Address of the Authorised Representative*
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* *for example:* *EMDN code(s)* *manufacturer list* *authorised representative list*
List of generic device group(s) covered by this mandate:	