






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

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