

2025-03-19 Version 1

List of Standard Fees for Conformity Assessment Activities under the MDR (2017/745)*, Notified body NB2862 Effective 1 January 2025

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	<u>Type of Fee</u>	<u>Fee</u> in local currency SEK	Factors influencing the calculation of fee charged	Fee range(mir max)
Administrative charges				
Application fee	N/A	None	No	N/A
 Administrative fee related to changes 	<u>Hourly</u>	3 000	Change Notice processing: Complexity of the change.	N/A
 Annual certificate maintenance fee (provide details which activities covered) 	<u>Flat</u>	36 190 - 69 780	For new MDR certificates an additional charge of 30 000 SEK will be added to the first annual fee. Factor influencing the calculation is number of employees at time of MDR application or beginning of the year.	36 190 - 69 780
• Other (specify)	N/A	N/A	N/A	N/A
European (Continent) Travel time costs (excluding expenses such as hotel costs)	<u>Flat</u>	Travel cost Up to 3 audit days 13 500 Travel cost		N/A
		3 days or more audit days 15 000		
Non-European (Continent) Travel time costs (excluding expenses such as hotel costs)	<u>Flat</u>	Travel cost Up to 3 audit days 65 000		
		Travel cost 3 days or more audit days 70 000		
Administrative costs related to handling of external services (laboratories, consultation or travel expenses)	N/A	N/A	No administrative costs related to handling of external services	N/A





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	Type of Fee	<u>Fee</u> in local currency SEK	Factors influencing the calculation of fee charged	Fee range(min- max)
Auditing				
 Audit (Certification; Recertification; Surveillance; Subcontractor/Supplier) 	<u>Daily</u>	41 000	Per audit day including audit planning, audit report, review of first response Corrective Action Plans, Independent Review and Certification Authority.	N/A
Unannounced Audit	<u>Flat</u>	105 000	Per unannounced audit	N/A
roduct testing				
 Laboratory testing (including preparation and reporting but excluding expenditures incurred for external tests) 	<u>Flat</u> <u>Hourly</u> <u>Daily</u>	TBD	To be determined and communicated to client before testing	TBD
ocumentation Review				
 Technical documentation assessment 	<u>Flat</u>	163 000	There are additions based on technologies, class and file complexity.	N/A
 Clinical evaluation report assessment (CEAR) 	<u>Hourly</u>	5 500	TD assessment include initial clinical assessment. Clinical oversight is charge by hour	N/A
 Expert panel consultation 	<u>Flat</u> <u>Hourly</u> <u>Daily</u>	TBD	To be determined and communicated to client before consultation	N/A
 Validation of the Summary of Safety and Clinical Performance (SSCP) 	<u>Hourly</u>	5 000	N/A	N/A
 Consultation with medicinal product authorities 	<u>Flat</u> <u>Hourly</u> <u>Daily</u>	TBD	To be determined and communicated to client before consultation	TBD
 Consultation with human tissue and cells competent authority 	<u>Flat</u> <u>Hourly</u> <u>Daily</u>	N/A	Not applicable out of scope	N/A
 Consultation with the coordinating competent authority for devices utilizing animal tissues 	<u>Flat</u> <u>Hourly</u> <u>Daily</u>	N/A	Out of scope, pending scope expansion with designating authority	N/A
 Evaluation/review of the Periodic Safety Update Report (PSUR) 	<u>Hourly</u>	4 000	N/A	N/A
Assessment of changes	<u>Flat</u> <u>Hourly</u> <u>Daily</u>	3000	Depending on the nature of change additional audit and file assessment maybe required to be determined	TBD





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			and communicated to client before assessment	
Reporting (if not covered above)	<u>Flat</u> <u>Hourly</u> <u>Daily</u>	N/A	Included in above charges	N/A
Special conditions for manufacturers belonging to SME as defined in Recommendation 2003/361/EC	Onsite audit dur	ation is calculated ba	ased on the number of full-tim	e employees.

* Article 50 of MDR (2017/745) & MDCG 2023-2 List of Standard fee

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