

List of Standard Fees According to (EU) 2017/745 Regulation

A) APPLICATION FEES

Product Risk Class	Fees	Туре	Remarks
Class Is, Im, Ir	1000 Euro		If more than one device is involved, the fee for highest product risk class is applied.
Class IIa	1500 Euro		
Class IIb (non-implantable)	2000 Euro		
Class IIb (implant)	2500 Euro	FLAT	
Class IIb Active (Administering removing medicines and other substances)	3000 Euro	- FLAT	
Class III	3500 Euro		
Class III implant	4000 Euro		

B) ANNUAL CERTIFICATE USAGE FEE

Product Risk Class	Fees	Туре	Remarks
Class Is, Im, Ir	5000 Euro		If more than one device is involved, the fee for highest product risk class is applied.
Class IIa	7000 Euro		
Class IIb (non-implantable)	9000 Euro		
Class IIb (implantable)	11000 Euro	FLAT	
Class IIb Active (Administering removing medicines and other substances)	13000 Euro	- FLAT	
Class III	15000 Euro		
Class III implant	17000 Euro		

C) AUDIT MAN/DAY FEE

Location	Fees	Туре	Remarks	
EU Countries	2500 Euro	DAILY (MAN/DAY)	Calculated mainly based on IAF MD-9 by applying several	
Others	2500 Euro		increasing and decreasing factors.	

1/3 FR.MED.165 R.05

List of Standard Fees According to (EU) 2017/745 Regulation

D) TECHNICAL DOCUMENTATION REVIEW MAN/DAY FEE

Location	Fees	Туре	Remarks
EU Countries	2500 Euro	DAILY (MAN/DAY)	Time spent will be calculated based on the product risk class. Following factors will increase the duration per device, - PSUR, PMCF, SSCP, PMS Reviews -Routine reviews on Technical Documentation changes - Devices in sterile condition and number of applied sterilization methods - Devices requiring biocompatibility review - Devices incorporating software
Others	2500 Euro		 Devices that are absorbable or locally dispersed Pre-market clinical investigation review Medicinal Product Authority Consultation Clinical Evaluation Consultation Procedure Consultation procedure for devices that are systemically absorbed

E) ADMINISTRATIVE AND OTHER FEES

Task	Fees	Туре	Remarks	
Initial review on changes	150 Euro	FLAT	Fees to be paid to the authorities are to be invoiced separately based on the rates available during the	
Administrative task for outgoing transfers	500 Euro	FLAT		
Preparation and follow up activities for authority consultations	100 Euro	HOURLY		
Assessment on appeals	100 Euro	HOURLY	consultation process.	
Travel Time (excluding travel and accommodation expenses)	50 Euro	HOURLY		

2/3 FR.MED.165 R.05



List of Standard Fees According to (EU) 2017/745 Regulation

F) FEES FOR REPEATING NON-CONFORMITY CORRECTIONS

Task	Fees	Туре	Remarks
Repeating Non-conformity Correction	350 Euro	FLAT	The contract will include one-time review of the non-conformities. For repeating non-conformity reviews within defined deadline will be invoiced separately. Repeating reviews will not be conducted once the deadlines are reached.

G) FEE CALCULATION PARAMETERS

Type of Assessment	Application Fee	Annual Certificate Usage Fee	Audit Fee	Technical Documentation Review Fee
Initial Assessment	+	+	+	+
Surveillance Assessment	-	+	+	+
Re-Assessment	+	+	+	+
Transfer Assessment (From another Notified Body to SZUTEST)	+	+	+	0
Transfer Assessment (From SZUTEST to another Notified Body)	**	-	-	-
Change Assessment	***	-	0	0
Scope Extension Assessment	+	*	0	0
Unannounced Site Audit	-	-	+	-
Follow Up Audit	-	-	+	-

^{&#}x27;o': Optional

H) SPECIAL CONDITIONS FOR MANUFACTURERS BELONGING TO SMEs AS DEFINED IN RECOMMENDATION 2003/361/EC

%3 of discount is applied for SMEs from the total initial and re-certification contract amount.

3 / 3 FR.MED.165 R.05

^{&#}x27;+': to be calculated

^{&#}x27;-': not to be calculated.

^{*} For higher product classes

^{**} The expenses to be invoiced for administrative work for controlling the completeness of transfer assessment submission is 150 EUR.

^{***} The expenses to be invoiced for administrative work in case of a change from SZUTEST to another notified body is 500 EUR.