

List of Standard Fees for Conformity Assessments Activities under the IVDR (2017/746)

Notified Body DEKRA Certification B.V. (NB 0344)

| | Type of Fee | Fee | Factors influencing the calculation of fee charged | Expected fee range (min-max). In special cases fees can differ |
|---|-------------|---------|---|--|
| Administrative charges | | | | |
| Application fee | Flat | € 1.800 | Complexity of the applications, number of applications | ≥ € 1.800 |
| Administrative fee related to changes | Hourly | € 450 | Complexity of the change | ≥ € 450 |
| Annual certificate maintenance fee | Flat | € 2.340 | First certificate - costs for regulatory requirements that apply to the accreditations as an auditing organization | € 2.340 |
| | | € 1.300 | Per additional certificate - costs for regulatory requirements that apply to the accreditations as an auditing organization | € 1.300 |
| Travel time costs | Hourly | € 162 | Location of manufacturer | € 162 - € 1.296 (per hour - day) |
| Administrative costs related to handling of external services (laboratories, consultation or travel expenses) | Hourly | € 450 | Testing or consultation by third party is billed at cost | ≥ € 450 |
| Auditing | | | | |
| Audit (Certification; Recertification; Surveillance; Subcontractor/Supplier) (including reporting and project management & excluding travel time and expenses). | hourly | € 306 | Number of FTE's, increase/decrease factors (MD9) | € 2.450 - € 4.900 (per 0,5 - 1 day) |
| Unannounced Audit | Daily | € 3.615 | Audit tariff per day, preparation and reporting | € 3.615 - € 7.230 |
| Product testing | | | | |
| Laboratory testing for verification of performance (including preparation and reporting but excluding expenditures incurred for external tests) | Hourly | € 450 | Dependent on complexity of the submitted product(s) | ≥ € 450, plus costs incurred for external testing will be invoiced at cost |
| Batch testing | Hourly | € 450 | Dependent on complexity of the submitted product(s) | ≥ € 450, plus costs incurred for external testing will be invoiced at cost |

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| Documentation review | | | | |
| Technical documentation assessment <u>Including</u> Project management and certification management | Hourly | € 450 | Dependent on risk classification/complexity of the submitted file(s). | € 3.600 - € 72.000 (1-20 days) |
| Performance Evaluation Assessment Report (PEAR) | Hourly | € 450 | Part of the technical documentation assessment and related project management | See Technical documentation assessment above |
| Expert panel consultation | Hourly | € 450 | Part of the technical documentation assessment and related project management | See Technical documentation assessment above |
| Validation of the Summary of Safety and Performance (SSP) | Hourly | € 450 | Dependent on complexity of the submitted file(s). Part of the PSUR report | See Evaluation/review of the Periodic Safety Update Report below |
| Consultation of a medicinal product authority for a companion diagnostic | Hourly | € 450 | Dependent on complexity of the submitted file(s) | ≥ € 450, plus costs incurred for external consultation will be invoiced at cost |
| Consultation of an EU reference laboratory for performance verification | Hourly | € 450 | Dependent on complexity of the submitted file(s) | ≥ € 450, plus costs incurred for external consultation will be invoiced at cost |
| Consultation of an EU reference laboratory for batch testing | Hourly | € 450 | Dependent on complexity of the submitted file(s) | ≥ € 450, plus costs incurred for external consultation will be invoiced at cost |
| Evaluation/review of the Periodic Safety Update Report (PSUR) | Hourly | € 450 | Dependent on complexity of the submitted file(s) | € 3.600 - € 7.200 (1 - 2 days) |
| Assessment of changes | Hourly | € 450 | Dependent on complexity of the change(s) | ≥ € 450 |
| Special conditions for manufacturers belonging to SME as defined in Recommendation 2003/361/EC 6 | SME manufacturers and first-time applicants benefit from reduced audit days based on FTEs, a 20% discount on application and a free onboarding package addressing Notified Body processes and regulatory aspects to enhance the conformity assessment process cost-efficiency. | | | |

Disclaimers:

Expected fee range might differ due to the complexity of the dossier(s). Reporting is covered in the tariffs shown above. Please note that the information provided is indicative and does not confer any rights. It is not tailored to your specific circumstances. For detailed information and an accurate quotation, we encourage you to contact us and request a formal proposal for our services.

For questions about our services, please contact medical.nl@dekra.com or call +31(0)88 96 83005.

Find out more about how DEKRA can support you by visiting [our website](#).