

2025 Biocidal Product Fees for Types 1 to 19 (TİTCK)

SERVICE CONTENT	2025 FEE (TRY)
Completeness check according to Articles 12 and 14 of the Biocidal Products Regulation (for each product type as per Annex V)	11.753,62
Completeness check according to Articles 12 and 14 of the Biocidal Products Regulation (including the verification of additional documents as per Annex IIIB)	14.653,24
Completeness check according to Articles 12 and 14 of the Biocidal Products Regulation for an additional product type (for each product type as per Annex V)	11.729,71
Evaluation fee for product authorization according to Articles 12 and 14 of the Biocidal Products Regulation for each product type	57.741,61
Product authorization evaluation fee according to Articles 12 and 14 of the Biocidal Products Regulation for each product type (including additional documents as per Annex IIIB)	80.541,95
Evaluation of product type addition/modification according to Articles 12 and 14 of the Biocidal Products Regulation	57.707,92
Evaluation for the addition of a harmful substance according to Articles 12 and 14 of the Biocidal Products Regulation	8.828,05
Completeness check for each product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation	3.695,47
Completeness check for each product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation (including the verification of additional documents as per Annex IIIB)	3.935,32
Completeness check for an additional/modification of a product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation	3.660,78
Registration and framework formulation evaluation fee for each product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation (including additional documents as per Annex IIIB)	29.307,10
Evaluation of product type addition according to Articles 13, 15, and 16 of the Biocidal Products Regulation for each product type	29.301,39
Evaluation for the addition of a harmful substance according to Articles 13, 15, and 16 of the Biocidal Products Regulation	4.433,60

2025 Fees for Biocidal Products (Excluding Types 1-19) (General Directorate of Public Health)

SERVICE CONTENT	2025 FEE (TRY)
Completeness check according to Articles 12 and 14 of the Biocidal Products Regulation (for each product type as per Annex V))	11.750
Completeness check according to Articles 12 and 14 of the Biocidal Products Regulation (including the verification of additional documents as per Annex IIIB	14.659
Completeness check according to Articles 12 and 14 of the Biocidal Products Regulation for an additional product type (for each product type as per Annex V	11.750
Evaluation fee for product authorization according to Articles 12 and 14 of the Biocidal Products Regulation for each product type	57.764
Product authorization evaluation fee according to Articles 12 and 14 of the Biocidal Products Regulation for each product type (including additional documents as per Annex IIIB)	80.553
Evaluation of product type addition/modification according to Articles 12 and 14 of the Biocidal Products Regulation	57.727
Evaluation for the addition of a harmful substance according to Articles 12 and 14 of the Biocidal Products Regulation	8.807
Completeness check for each product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation	3.673
Completeness check for each product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation (including the verification of additional documents as per Annex IIIB)	3.935
Completeness check for an additional/modification of a product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation	3.673
Registration and framework formulation evaluation fee for each product type according to Articles 13, 15, and 16 of the Biocidal Products Regulation (including additional documents as per Annex IIIB)	29.321
Evaluation of product type addition according to Articles 13, 15, and 16 of the Biocidal Products Regulation for each product type	29.321
Evaluation for the addition of a harmful substance according to Articles 13, 15, and 16 of the Biocidal Products Regulation	4.458

Review and dispatch of application along with records and documents (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	11.729,71	Review and dispatch of application along with records and documents (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation))	11.750
Completeness check and evaluation of additional data as per Annex IIIA of the Biocidal Products Regulation (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB)	117.210,62	Completeness check and evaluation of additional data as per Annex IIIA of the Biocidal Products Regulation (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB)	117.190
Completeness check (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	96.735,75	Completeness check (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	96.720
Evaluation of data and documents, including the assessment of additional data as per Annex IIIA of the Biocidal Products Regulation (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB)	849.816,37	Evaluation of data and documents, including the assessment of additional data as per Annex IIIA of the Biocidal Products Regulation (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB)	849.811
Evaluation of data and documents for acceptance, in accordance with the Biocidal Products Regulation (for the inclusion of a new active substance in Annex I, Annex IA, or Annex IB)	762,66	"Issuance of Product Notification and Registration Certificate for Biocidal Products without Active Substances, etc.	22.137
Review and dispatch of application along with records and documents (for the inclusion of a microorganism as an active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	11.022,98	Evaluation Fee for Each Product Included in the Product Notification and Registration Certificate for Biocidal Products without Active Substances, etc., and for Each Additional Product to be Included	1.134
Completeness check (for the inclusion of a microorganism as an active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	73.672,92	Biocidal Products without Active Substances, etc. Product Document Renewal Fee (for additional product inclusion, loss, etc.)	4.422
Evaluation of data and documents for acceptance (for the inclusion of a microorganism as an active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	512.812,33	Biocidal Products without Active Substances, etc. Product Label Change Evaluation Fee (per product)	762
Product label registration for biocidal products (per product)	22.583,38	Evaluation of Data and Documents for Acceptance (as per the Biocidal Products Regulation, for the addition of a new active substance to Annex I, Annex IA, or Annex IB)	759.512
Biocidal product health and free sale certificate fee (per product)	1.283,14	Review and dispatch of application along with records and documents (for the inclusion of a microorganism as an active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	11.039

Biocidal product authorization transfer fee (per product - import/production)	65.697,14
Biocidal product production site change/production site addition/packaging site addition fee (per product)	32.232,67
Biocidal product name change/company name change fee (per product)	32.232,67
Biocidal product document renewal fee (per product)	3.695,47
Biocidal product authorization fee	110.350,68
Biocidal product framework formulation authorization fee (per product)	29.307,10
Biocidal product registration fee (per product)	43.944,45
Biocidal Product Applicator Certificate Fee	137,29

Completeness check (for the inclusion of a microorganism as an active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	73.674
Evaluation of data and documents for acceptance (for the inclusion of a microorganism as an active substance in Annex I, Annex IA, or Annex IB of the Biocidal Products Regulation)	512.812
Product label registration for biocidal products (per product)	22.137
Biocidal Products Responsible Manager Training; Training and Course Registration Fee (42 hours)	22.137
Biocidal Products Responsible Person Update Training Fee (8 hours)	4.271
Biocidal Product Analysis Authorization Certificate Fee	65.213
Biocidal Product Analysis Authorization Certificate Renewal Fee	26.074
Fee for the Biocidal Product Application Training Organization Authorization Certificate issued to relevant professional organizations and universities wishing to provide applicator training (excluding Provincial Health Directorates)	13.851
Biocidal Product Application Training Organization Authorization Certificate Renewal Fee for relevant professional organizations and universities wishing to provide applicator training (excluding Provincial Health Directorates)	3.462
Biocidal Product Framework Formulation Completeness Check Fee (separately for each framework formulation)	3.673
Biocidal Product Framework Formulation Evaluation Fee (separately for each framework formulation)	29.321
Active Substance Supplier Change Evaluation Fee	8.807