

1st Annex F1A/15 to the Certificate No 244-7

SCOPE of ACCREDITATION

of the

Laboratory for Chemical Analysis of Medicines of the

Division of Laboratories, National Organisation for Medicines

Types of test/Properties	Applied standards/Techniques		
Chemical tests			
According to the monographs of current editions of European Pharmacopoeia (EP), Hellenic Pharmacopoeia (ΕΦ), British Pharmacopoeia (BP), United States Pharmacopoeia (USP) and approved validated methods of product files			
Identification and assay of active substance Test for related substances of active substance	High Performance Liquid Chromatography (HPLC) – (DAD/PDA) (MEE 9001)		
Identification and assay of active substance	Spectrophotometry UV-Vis (MEE 9003)		
Identification of active substance Test for related substances of active substance	Thin Layer Chromatography (TLC) (MEE 9005)		
Assay of active substance	Titrimetry (use of indicator) (MEE 9006)		
Potentiometric determination of pH	General method 2.2.3 EP of (MEE 9008)		
	Chemical tests raphs of current editions of European Pharmacoocia (BP), United States Pharmacopoeia (USP) Identification and assay of active substance Test for related substances of active substance Identification and assay of active substance Identification of active substance Test for related substances of active substance Test for related substances of active substance Assay of active substance		

Tested materials/products	Types of test/Properties	Applied standards/Techniques	
Single-dose pharmaceutical products	Test for uniformity of content	General method 2.9.6 EP using high performance liquid chromatography (HPLC) – (DAD/PDA)	
	Semi-micro determination of water	General method 2.5.12 EP (MEE 9011)	
	Physical tests		
According to the monographs of current editions of European Pharmacopoeia (EP), Hellenic Pharmacopoeia (ΕΦ), British Pharmacopoeia (BP), United States Pharmacopoeia (USP) and approved validated methods of product files			
Single-dose pharmaceutical products	Test for uniformity of mass / Test for average mass	General method 2.9.5 EP by weighing / specification of the product dossier	
Tablets and capsules	Test for dissolution	General method 2.9.3 EP and assay of active substance with HPLC-(UV-Vis) or Spectrophotometry UV-Vis	
		(MEE 9002)	
	Test for disintegration	General method 2.9.1 EP	
		(MEE 9010)	

Site of assessment: Permanent Laboratory premises, Mesogeion 284, Holargos, Athens Approved signatories: V. Violakis, E. Kyriakopoulou, E.Barmpagianni, P.Karava

This scope of Accreditation replaces the previous one dated 16.02.2022. The Accreditation Certificate No **244-**7, to ELOT EN ISO/IEC 17025:2017, is valid until 07.03.2026.

Athens, 6.05.2022



2nd Annex F1B/15 to the Certificate No 244-7

SCOPE of ACCREDITATION

of the

Microbiological Laboratory

of the

Division of Laboratories, National Organization for Medicines

Tested materials/products	Types of test/Properties	Applied standards/Techniques	
Microbiological tests			
Sterile pharmaceutical	Sterility tests	General methods PH.EUR 2.6.1, USP/NF <71> (MEE 7001)	
products and medical devices	Test for bacterial endotoxins	General methods PH.EUR 2.6.14, Method A (gel- clot: limit test) and Method B (gel-clot: semi- quantitative test) USP/NF <85> and <161> (MEE 7006)	
Non sterile pharmaceutical products and substances for pharmaceutical use	Test for microbial quality	General methods PH.EUR 2.6.12 and 2.6.13 (MEE 7002)	
Foodstuffs intended for particular nutrition	Detection of Salmonella spp	EN ISO 6579-1:2017/ Amd 1:2020 excluding Annex D (serovars Typhi and Paratyphi)	
S'A S	Detection of Cronobacter spp	EN ISO 22964:2017	

Site of assessement: Permanent Laboratory premises, Mesogeion 284, Holargos, Athens Approved signatories: V. Violakis, G. Bouziou, V. Birtsas.

This scope of Accreditation replaces the previous one dated 16.02.2022. The Accreditation Certificate No **244-**7, to ELOT EN ISO/IEC 17025:2017, is valid until 07.03.2026.

Athens, 6.05.2022



4th Annex F1D/15 to the Certificate No 244-7

SCOPE of ACCREDITATION

of the

Laboratory for Chemical Analysis of Feeding Stuffs, Dietary Products, Herbal Medicines

of the

Division of Laboratories, National Organization for Medicines

Tested materials/products	Types of test/Properties	Applied standards/Techniques	
Chemical tests			
According to the monographs of current editions of European Pharmacopoeia (EP), Hellenic Pharmacopoeia (ΕΦ), British Pharmacopoeia (BP) and United States Pharmacopoeia (USP) and approved validated methods of product files			
Pharmaceutical starting materials used in veterinary medicinal products and veterinary medicinal products	Identification and Assay of active substance	Spectrophotometry UV-Vis (MEE 9003)	
	Assay of active substance	Titrimetry (use of indicator) (MEE 9006)	
	Identification and Assay of active substance	High Performance Liquid Chromatography (HPLC) – (DAD/PDA)	
Physical tests			
According to the monographs of current editions of European Pharmacopoeia (EP), Hellenic Pharmacopoeia (ΕΦ), British Pharmacopoeia (BP) and United States Pharmacopoeia (USP) and approved validated methods of product files			
Single-dose veterinary medicinal products	Test of uniformity of mass	General method 2.9.5 EP by weighing (MEE 9009)	

Site of assessment: Permanent Laboratory premises, Mesogeion 284, Holargos, ATHENS Approved signatories: V. Violakis, K.Fakinou.

This scope of Accreditation replaces the previous one dated 16.02.2022.

The Accreditation Certificate No 244-7, to ELOT EN ISO/IEC 17025:2017, is valid until 07.03.2026.

Athens, 6.05.2022



5th Annex F1E/14 to the Certificate No **244**-7 **SCOPE of ACCREDITATION**

of the

Biological and Toxicological Laboratory

of the

Division of Laboratories, National Organisation for Medicines

Tested materials/products	Types of test/Properties	Applied standards/Techniques		
Chemical tests				
	According to the monographs of current editions of Hellenic Pharmacopoeia (ΕΦ), British Pharmacopoeia (BP), United States Pharmacopoeia (USP)			
Pharmaceutical starting materials and pharmaceutical products	Identification and assay of active substance	Spectrophotometry UV-Vis (MEE 9003)		
	Test for related substances of active substance	Thin Layer Chromatography (TLC) (MEE 9005)		
Physical tests				
According to the monographs of current editions of Hellenic Pharmacopoeia (ΕΦ), British Pharmacopoeia (BP), United States Pharmacopoeia (USP)				
Single-dose pharmaceutical products	Test for uniformity of mass / Test for average mass	General method 2.9.5 EP by weighing (MEE 9009) / specification of the product dossier		
Tablets and capsules	Test for dissolution	General method 2.9.3 EP and assay of active substance with HPLC-(DAD/PDA) or Spectrophotometry UV-Vis (MEE 9002)		
Tablets and capsules	Test for disintegration	General method 2.9.1 EP (MEE 9010)		

Site of assessment: Permanent Laboratory premises, Mesogeion 284, Holargos, Athens Approved signatories: V. Violakis, G. Volikakis. P. Nikolaou.

This scope of Accreditation replaces the previous one dated 16.02.2022. The Accreditation Certificate No **244-7**, to ELOT EN ISO/IEC 17025:2017, is valid until 07.03.2026.

Athens, 6.05.2022



6th Annex F1F/15 to the Certificate No 244-7

SCOPE of ACCREDITATION

of the

Laboratory for Chemical Analysis of Cosmetics and Other Products of the

Division of Laboratories, National Organisation for Medicines

Tested materials/products	Types of test/Properties	Applied standards/Techniques	
Chemical tests			
Cosmetic products	Identification and determination of 2-phenoxy-ethanol, methyl-, propyl-, and butyl-4-hydroxybenzoate (parabens)	Annex /7th Commision directive 96/45/EC High performance liquid chromatographic (HPLC)-(DAD/PDA) (MEE 6005) Annex/ 6th directive 95/32 EC	
	Identification and determination of benzoic acid, 4-hydroxybenzoic acid, salicylic acid, sorbic acid	High performance liquid chromatographic (HPLC)-(DAD/PDA) (MEE 6006)	
Physical tests			
Medical gloves for single use	Requirements and testing for freedom from holes	ELOT EN 455-1:2020 (MEE 6011)	
Natural rubber latex male condoms	Determination of bursting volume and pressure	ISO 4074:2015 (MEE 6003)	

Site of assessment: Permanent Laboratory premises, Mesogeion 284, Holargos, Athens Approved signatories: V. Violakis, K. Venetsanou.

This scope of Accreditation replaces the previous one dated 16.02.2022. The Accreditation Certificate No **244-**7, to ELOT EN ISO/IEC 17025:2017, is valid until 07.03.2026.

Athens, 6.05.2022