



GELATIN CAPSULE

CERTIFICATE OF ANALYSIS

CUSTOMER: OMFE S.A.
ITEM DESCRIPTION: 2AJAAJA QG
CUSTOMER ITEM: 30200101
LOT NUMBER: R2500897
LOT QUANTITY APPROVED: 1910000
PRINTING: NOT APPLICABLE
INK COLOR: NOT APPLICABLE

COUNTRY: Spain
SIZE: 2
CUSTOMER ORDER: 4900064340-2
NAME: 2 BLANCO
APPROVAL DATE: 17/04/2025
MANUFACTURING DATE: 30/03/2025
EXPIRY DATE: 5 YEARS

FINISHED PRODUCT TESTS

PARAMETERS	LIMITS	REFERENCE	RESULT
MICROBIOLOGICAL AND CHEMICAL TEST			
Total Aerobic Microbial Count (TAMC)	10 ³ CFU/G	EP/USP	≤ 10
Salmonella	Negative / 10 G	EP/USP	Meets test
E. Coli	Negative / 1 G	EP/USP	Meets test
S. Aureus	Negative / 1 G	EP/USP	Meets test
Ps. Aeruginosa	Negative / 1 G	EP/USP	Meets test
DIMENSIONAL AND PHYSICAL TEST			
Diameter	Cap (mm)	Meets spec	TM*
	Body (mm)	Meets spec	TM*
Length	Cap (mm)	Meets spec	TM*
	Body (mm)	Meets spec	TM*
End thickness	Cap (micr)	Meets spec	QS*
	Body (micr)	Meets spec	QS*
Weight (mg)	Meets spec	TM*	63,2 (Average)
Moisture (%)	Meets spec	TM*	14,9 (Average)

	VISUAL	ATTRIBUTES			
VISUAL DEFECTS	Critical	Major	Minor	TM*	Meets
AQL (%)	0,010	0,040	0,25		
PRINTING DEFECTS	Critical	Major	Minor	TM*	Meets (if applicable)
AQL (%)	0,010	0,040	1,0		

*TM = Qualicaps Technical Manual *QS = Qualicaps internal specification "Meets test" means conforms to specification

METAPHARMACEUTICAL

N DE LOTE:

0230425

Signature 28/04/2023

Page 1 of 2

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CAPSULE COLOR FORMULATION

The design weight of a size 2 capsules is 63,0 mg , the cap representing 40% and the body 60% of the capsule weight. The mean capsule weight of 100 capsules can vary within the range of 58,3 - 67,7 mg.

CAP :

CODE	TITLE	COMPONENT PER CAPSULE (g/100g)
AJA	OPAQUE WHITE	TITANIUM DIOXIDE 2.000 WATER (14,5%) TARGET MOISTURE GELATIN

BODY :

CODE	TITLE	COMPONENT PER CAPSULE (g/100g)
AJA	OPAQUE WHITE	TITANIUM DIOXIDE 2.000 WATER (14,5%) TARGET MOISTURE GELATIN

COMPONENT	C.I. NUMBER	E.E.C. NUMBER
TITANIUM DIOXIDE	77891	E171

REGULATORY COMPLIANCE

RAW MATERIAL

GELATIN: Complies with the requirements of current EP, JP and USP/NF editions.

It is of pure bovine origin, complies with revision in force of European Guideline EMEA/410/01. Each supplier has a Ph. Eur certificate of suitability for their product (R1-CEP-2000-027-Rev.02, R1-CEP-2000-029-Rev.05, R1-CEP-2000-029-Rev.06, R1-CEP-2000-045-Rev.04, R1-CEP-2000-045-Rev.06, R1-CEP-2002-110-Rev.00, R1-CEP-2000-050-Rev.02).

COLORANTS: Comply with Directive 2009/35, Commission Regulation 231/2012 and where applicable, with the requirements of the EP and USP/NF.

PRINTING INKS: Comply with pharmaceutical regulations.

CAPSULES:

Do not contain preservatives and have not been treated with Ethylene Oxide. They comply with The European Agency for the Evaluation of Medicinal Products (EMA) guideline CPMP/ICH/283/95 and the European Pharmacopoeia for residual solvents.

DISINTEGRATION: Less than 15 minutes by the Ph. Eur. test and have a rupture time of less than 5 minutes by the USA Federal Specification 285A Acid Solubility Test.

This is to certify that the information above has been approved by QA as conforming to Qualicaps Europe specifications, as described in the current technical manual and applicable regulatory requirements.

ADRIANA OPREA
QA/QC MANAGER



Page 2 of 2

DATE: 25/04/2025

