ENTEGRIS CLEANING PROCESS
E C P
An Entegris Company

CONTROL REPORT

Customer	Management Quality Manual n° MMMQ0000 (contractual)
JOHNSON Diversey LTD	

Product : Clearklens IPA VH1 70%	**Control report n° :** CERT 9629B
Specification reference : NA	
Analyst : FGA	**Purchase order n° :** 4700732791
	Date of receipt : 10/04/2009

CONTRÔLE DES PROPRIETES PHYSIQUES ET CHIMIQUES

Customer's batch n° : ENT06291 09 100	**ISOTRON certificate n° :** 14062101
Date of manufacturing : 10/04/2009	**Produced quantity :** 167 x 5L
Expiery date : 04/2011	**FIDT N° :** FIDT 1814 D
ECP's batch n° : 06291	**Date of analysis :** 10/04/09

Résultats :

Analyzed characteristics	Method of analysis	State in the production	Specification
		Beginning	
Appearance 20°C	NA	C	Clear, colourless liquid
Specific gravity (20°C)	IDT0301	0,870	0,865 - 0,875
pH (20°C)	NA	9,0	6,0 - 9,0
Microbiological contagion of the EDI	IDT0236	< 1 UFC/100mL	< 10 cfu/100mL

C : Conform NC : Not Conform
NA : Not Applicable

Conclusion : [X] CONFORM [ ] NOT CONFORM

Date : 30/06/09	Edited: L. GRESSE	Checked: D. CHEUNG

CERTIFICATE OF TREATMENT BY GAMMA RADIATION

WE UNDERSIGNED :

Isotron France S.A.S.
MIN 712
13323 MARSEILLE CEDEX 14 - FRANCE

ECP Reçu le
29 MAI 2009
N° 090529 JLA

CERTIFY THAT WE TREATED BY GAMMA RADIATION ACCORDING TO THE SPECIFIC CUSTOMER'S REQUIREMENTS AND TO :

- the specification of treatment # 224503P
- The requirements of the European Pharmacopoeia
- the results of the dosemapping # 110505 of 30/03/2006

THE FOLLOWING PRODUCTS : (according to the customer's indication)

Customer	:	**ECP ENTEGRIS CLEANING PROCESS**
Product	:	5L BOTTLES KITS
Customer's reference	:	ORDER # CF 090074 OF 02/04/09
Quantity	:	1 PALLET
Irradiation date	:	2009.04.05
Irradiation dose	:	16.5 kGy to 21.4 kGy
Irradiation batch number	:	14062101

Clariflens IPA VHJ 70%
Lot ENT 062J1 09 J00
Date fab : J0/04/09
Date pér : 04/20 M
J. BURGOS le 29/06/09

The control of the applied radiation dose is done by Isotron France SAS using Red Perspex dosimeters calibrated by the English National Physical Laboratory.

Isotron France S.A.S.,

S. DEFAZIO
Process Control

C. SIMONIN
Quality Officer

Certificate # 14062101 / 1

English translation of the french original certificate.

charles river
accelerating drug development. exactly.

RAPPORT D'ESSAI / *ANALYSIS REPORT*

N° 60-136-1 – Page / *Page* 1 / 1

22 AVR. 2009

Edité le / *Printed on the* : 20 avril 2009, par / *by* :

pour le client / *for customer* n° : 7669

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ECP (Entegris Cleaning Process)
A l'attention Mr PORTEFAIX Jérôme
395 rue Louis Lépine
34000 MONTPELLIER

ESSAI DES ENDOTOXINES BACTERIENNES
Bacterial Endotoxins Test

Contrat Technique
Technical Contract

Questionnaire technique Selon 58-036

Technique / *Method*

Méthode D
Colorimétrie cinétique

Sensibilité / *Sensitivity*

0,005 UI/mL

Date de réception
Simple delivery date

16 avril 2009

Date d'analyse
Testing date

17 avril 2009

Nbre d'échantillons
Number of samples

1

Produit / *Product*
ClearKlens

Limite en endotoxines / *Endotoxin limit* (Entegris)
0,25 UI/ml

Les essais ont été réalisés en accord avec la Pharmacopée Européenne en vigueur, chapitre "2.6.14 – Endotoxines Bactériennes" harmonisé avec les Pharmacopées Américaine et Japonaise.
The assay was performed in compliance with the European Pharmacopoeia in force, chapter "2.6.14 – Bacterial endotoxins", harmonized in collaboration with the American and Japanese Pharmacopoeias.

ID produit / *product ID*
ClearKlens IPA 70% 5L

Lot / Prélèvement / Numéro *Batch / Sample / Number*	Concentration en endotoxines *Endotoxin amount*	Unités *Units*
ENT06291 09 100	**< 0,25**	**UI/mL**

cofrac
ESSAIS
ACCREDITATION
N° 1-1850
Scope available
on www.cofrac.fr

Opérateur / *Operator*
Fanny MORI
Tech. de laboratoire / *Lab. Technician*
20 AVR. 2009

Approbateur Qualité / *Quality Reviewer*
N.A

Approbateur Technique / *Technical Reviewer*
Gilles GOY
Responsable Technique Laboratoire
20 AVR. 2009

ISO 9001:2000 SGS

MedicalLab

Report n° :	09/OI/STE/060
Sponsor :	ECP 395, rue Louis Lépine 34000 MONTPELLIER

Test report Essay of Sterility - Method filtration on membranes

According to the protocol 2.6.1 described into the sixth Europeen Pharmacopee Edition

TEST ARTICLE

Name of product :	ClearKlens IPA VH1	Order number :	CF090335
Customer reference :	7516393	Internal reference :	-
Batch number :	ENT 06 291 09 100	Material :	-
Date of receipt :	16 avril 2009	Data sterilization :	-
Date of test :	2 juin 2009	Comment(s) :	3/4 de production 1/4 de production
Quantity of used sample :	2		

PROTOCOL

Tested volume of product : 500 ml

Neutralizing solution : DNP + Thiosulfate

Rinsing volume : 3 x 100 ml

Number of media tested : 2

Immersion volume of membranes : 90 ml

Conditions	Media	Incubation temperature	Time of incubation
Aerobic and fugal	Tryptone soy solution	20 ± 5°C	14 days
Anaerobic and aerobic	Thioglycolate Résazurine solution	30 ± 5°C	14 days

Fertility validation : 09/OI/VAL.FER/006

RESULTS

Conditions	Assessment of the media turbidimetry			
	After 7 days		After 14 days	
Tryptone soy solution	0	positive	0	positive
	1	negative	1	negative
Thioglycolate Résazurine solution	0	positive	0	positive
	1	negative	1	negative

CONTROLS

Work plan control (before and after) :	0	0
Air control :	0	

Glove control :	Conforme
Media sterility control :	Conforme

CONCLUSION

The tested product doesn't shown any microbian development after 14 days of incubation.

No *product has been positive during the test.*

Redact by : PEGOUD Céline, Technicien Biologiste

Approved by : MARTINHO Alice, Ingénieur Biologiste

Date : vendredi 26 juin 2009

Results and conclusion apply only on the test articie tested. Any extrapolation of these data to other samples is the responsibility of the Sponsor.

MedicalLab
Microbiological and physico-chemical analysis - process validation
EN ISO 13485 (2003)

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