GELDA SCIENTIFIC

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ISO 17025
MOE -
LICENCE #2203

LABORATORY ANALYSIS REPORT

Phoenix Biomedical Products S.L

REPORT NO.:	**594**	**Date Reported:**	**30-Sep-16**
Date Received:	12-Sep-16	Analysis Date:	14-Sep-16
Smpl.Code From:	1	Fax:	NA
Smpl.Code To:	35	Tel:	NA
# of Samples:	35	Invoice Address:	c/Verde, Parcela 3.7
Date Released:	30-Sep-16		Parque Tecnológico Fuente Álamo
Mail Reports To:	**PHOENIX BIOMEDICAL PRODUCTS INC.** 14-30 Eglinton Avenue West, Suite 273 Mississauga, ON L5R 0C1		Ctra. del Estrecho-Lobosillo Km.2 30320 Fuente Álamo, Murcia, SPAIN
		Email:	stardish@phoenix-biomed.com

Certificate of Sterility

Product: Petri Dishes

Lot No.s

G19180	H01057	H30110
G26110	H02360	H31360
G26360	H02057	H31110
G27360	H03360	I01360
G27110	H04057	I02360
G28360	H04360	I05110
G28110	H05057	I05360
G29110	H05360	I06110
G29360	H09057	I06360
G30057	H29360	I07110
G30360	H29110	I07057
H01360	H30360	

Test Method: Modified USP <71>, Sterility Tests – Phoenix testing protocol (SOP#:GL1005-Not ISO 17025 nor Health Canada accredited)

Number of Units Tested per lot #: 4

Test Results:

	Fluid Thioglycollate Agar					Soybean-Casein Digest Agar				
Incubation Time	14 days /32.5°C +/- 2.5°C					14 days /22.5°C +/- 2.5°C				
Test Product	1	2	3	4		1	2	3	4	
	NG	NG	NG	NG		NG	NG	NG	NG	
Negative Control: (Medium only) [TSBA #1905525, FTGA #1865823]	NG					NG				

Legend: NG- No Growth; G: Growth

Authorized By:
Dr.Sumona Guha
Quality/Lab Manager

Verified by
Mr. Bhavdeep Chandhoke
Lab Supervisor