

840 RESEARCH PARKWAY, SUITE 546 OKLAHOMA CITY, OK 73104 PHONE (405) 271-1144 FAX (405) 271-1174

Certificate Of Analysis

CLIENT: BET Pharmacy-KY

Web Report

ARL #: 450801-01 LOT #: 29332@2

DESCRIPTION: P4LA Progesterone 150 mg/mL

DATE RECEIVED: 02/13/2018

STORAGE: 20°C to 25°C (68°F to 77°F)

CONTAINER: Twelve 100 mL amber vials w/ 100 mL each

Analyte / Specifications	Expected Amount	Units	Results	% Of EXP.	Test Method	Date Tested
Progesterone	150	mg/mL	151.1202	100.7%	HPLC	2/14/2018
Specifications = 90.0% - 110.0%						

Formulation ID: 06

The potency method(s) used for testing passed system suitability requirements per ARL SOP AMP-012 for non-cGMP analysis. Product specific method validation is not available for this sample and specification(s) are for informational purposes only. Client should verify the specification and analyte reported are correct for the compounded formulation.

Port	02/16/2018	
Ryan Kostuck - Laboratory Supervisor	Date Reported	ARL Form QUF-078-V4 03/05/2010



840 RESEARCH PARKWAY, SUITE 546 OKLAHOMA CITY, OK 73104 PHONE (405) 271-1144 FAX (405) 271-1174

Microbiology Report

CLIENT: BET Pharmacy-KY

Web Report

ARL #: 450801-01 LOT #: 29332@2

DESCRIPTION: P4LA Progesterone 150 mg/mL

DATE RECEIVED: 02/13/2018

STORAGE: 20°C to 25°C (68°F to 77°F)

CONTAINER: Twelve 100 mL amber vials w/ 100 mL each

ANALYSIS	Limits	Results	Test Method	Date Tested
Sterility (*Preliminary*)	Sterile / Not Sterile	No Growth at 3 Days	USP <71>	02/13/2018
Endotoxin	555.56 EU/mg	<0.07 EU/mg	USP <85>	02/16/2018
Fungal	Sterile / Not Sterile	No Growth at 6 Days	MBI-114	02/13/2018

Formulation ID: 06

The citation to USP <71> is conditioned on the accuracy of our customer's verification that the sampling program complies with the provisions of USP <71>.

Sterility – This preliminary report was issued after approximately 72 hours of incubation. In accordance with the test methodology, the sample will be incubated for 14 days; if there is any change in the sample a supplemental report will be issued.

Fungal – This preliminary report was issued after approximately 4 days of incubation. In accordance with the test methodology, the sample will be incubated for 14 days; if there is any change in the sample a supplemental report will be issued.

 $Endotoxin - To \ calculate \ the \ endotoxin \ limit \ use \ the \ following \ formulae: \ EL = K/M \ where \ K = tolerance \ limit \ (EU/kg) \ and \ M = Maximum \ dose/kg/hour \ or \ Maximum \ dose/kg$

ARL Form QUF-078-V6 11/26/2012

Parenteral: K is 5 EU/kg for any route of administration /Intrathecal: K is 0.2 EU/kg body weight)

Radiopharmaceutical parenteral: K is 175/V or Intrathecal radiopharmaceuticals: K is 14/V, where V is the maximum recommended dose in mL. Dermal Application: K/M, where K = 5 EU/kg and M is the (maximum dose/m2/hour \times 1.80 m2)/70 Kg.

Mern Hust Date Reported

Description 102/19/2018

Date Reported

Results reported above relate only to the sample that was tested.



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CONTAINER: Twelve 100 mL amber vials w/ 100 mL each

ANALYSIS	Limits	Results	Test Method	Date Tested
Sterility	Sterile / Not Sterile	No Growth at 14 Days (subcultured) *	USP <71>	02/13/2018
Fungal	Sterile / Not Sterile	No Growth at 14 Days (subcultured) *	MBI-114	02/13/2018

Formulation ID: 06

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* This is not a final result. After 14 days of incubation, the material being tested rendered the media turbid so that the presence or absence of microbial growth cannot be readily determined by visual examination. Per USP <71>, the sample will be inoculated into new growth media and incubated for an additional 4 days before the test is complete. An updated report will be issued at the conclusion of the test.

Kerri Hirst - Microbiologist II

Hern Hust

02/27/2018

Date Reported

Sterility - 14 day sterility report. In accordance with the test methodology, the sample was incubated for 14 days.

Fungal - 14 day fungal report. In accordance with the test methodology, the sample was incubated for 14 days.



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Microbiology Report

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DATE RECEIVED: 02/13/2018

STORAGE: 20°C to 25°C (68°F to 77°F)

CONTAINER: Twelve 100 mL amber vials w/ 100 mL each

ANALYSIS	Limits	Results	Test Method	Date Tested
Sterility	Sterile / Not Sterile	Sterile	USP <71>	02/13/2018
Fungal	Sterile / Not Sterile	Sterile	MBI-114	02/13/2018

Formulation ID: 06

The citation to USP <71> is conditioned on the accuracy of our customer's verification that the sampling program complies with the provisions of USP <71>.

Kerri Hirst - Microbiologist II

Hern Herst

03/05/2018

Date Reported

Sterility – 18 day sterility report. In accordance with the test methodology, the sample was incubated for a minimum of 18 days.

Fungal – 18 day fungal report. In accordance with the test methodology, the sample was incubated for a minimum of 18 days.