



U.S. Food and Drug Administration  
Office of Regulatory Affairs  
12120 Parklawn Dr.  
Rockville, MD 20852  
[www.fda.gov](http://www.fda.gov)

September 24, 2018

Mr. Rodney Rogan, General Manager  
Lacsa (PTY) Limited  
72 Ballantrae Road, Durban South,  
Kwazulu-Natal, 4052 South Africa

Dear Mr. Rogan:

The U.S. Food and Drug Administration (FDA) conducted an inspection at Lacsa (PTY) Limited, FEI 3003699242, located at 72 Ballantrae Road, Durban South, Kwazulu-Natal, 4052 South Africa, from 7/9/18 to 7/12/18. FDA has determined that the inspection classification of this facility is “no action indicated (“NAI”).<sup>1</sup> Based on this inspection, this facility is considered to be in an acceptable state of compliance with regards to current good manufacturing practice (CGMP).

This letter is not intended as an endorsement or certification of the facility. It remains your responsibility to ensure continued compliance with CGMP.

An inspection classification of NAI for CGMP compliance will not directly negatively impact FDA’s assessment of any pending marketing application referencing this facility. Please note, however, that application approval will depend on a product- and application-specific facility assessment conducted by CDER’s Office of Pharmaceutical Quality. This letter does not address or reflect FDA’s decision making with respect to any potential non-CGMP compliance issues.

FDA has concluded that this inspection is “closed” under 21 CFR 20.64(d)(3), and we are enclosing a copy of the narrative portion of the Establishment Inspection Report (EIR). It may reflect redactions made by FDA in accordance with the Freedom of Information Act (FOIA) and 21 CFR part 20. This, however, does not preclude you from requesting additional information under FOIA.

If you have any questions regarding this letter, you may contact Atul Agrawal via telephone at 240-402-4900 or email at [atul.agrawal@fda.hhs.gov](mailto:atul.agrawal@fda.hhs.gov).

Sincerely,

Kevin Gonzalez  
Branch Chief  
Foreign Pharmaceutical Quality Inspections

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<sup>1</sup> See Inspection Classification Definitions, at <https://www.fda.gov/ICECI/Inspections/ucm223231.htm>