



## Certificate of Good Manufacturing Practices

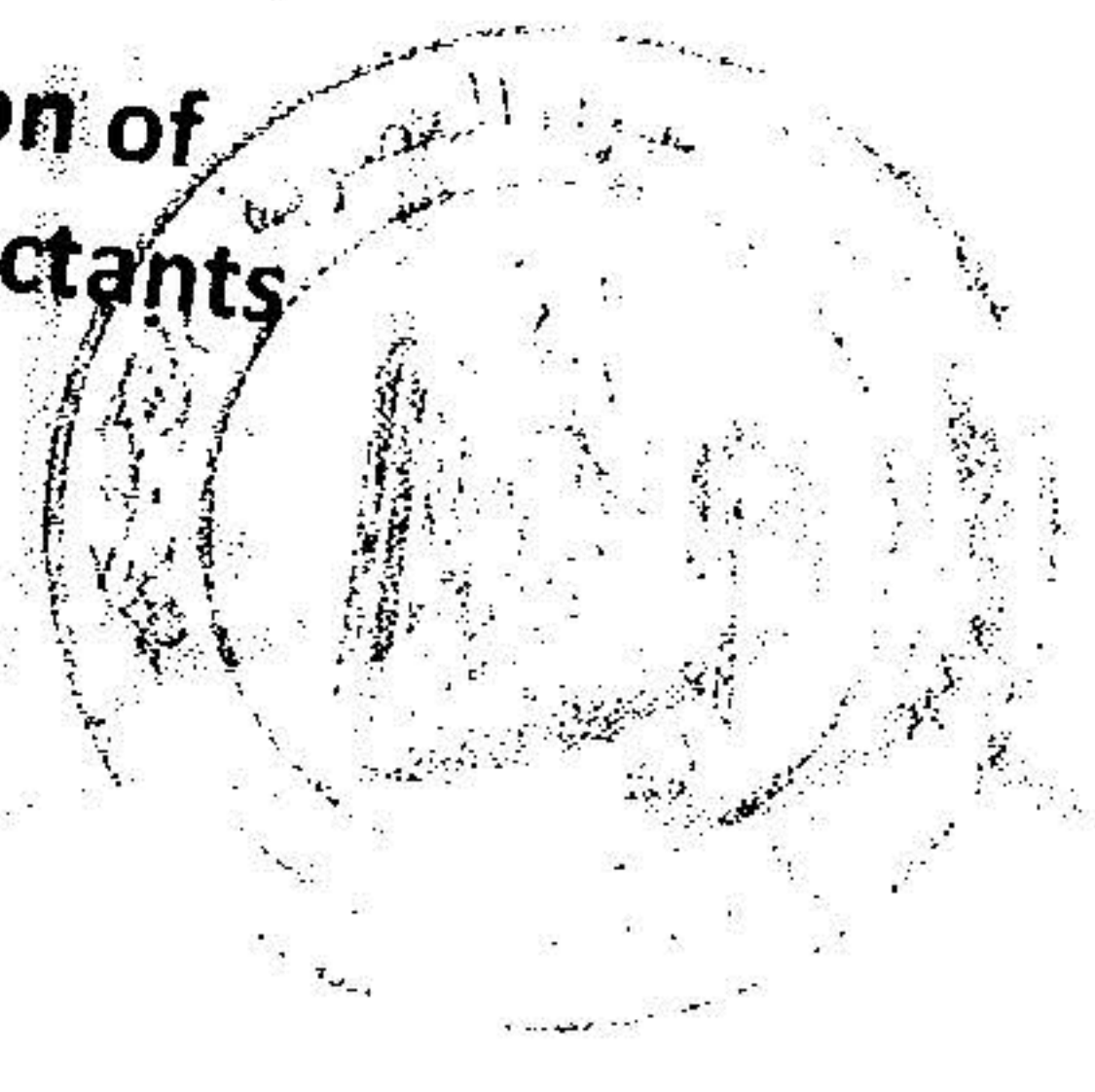
This certificate conforms to the format recommended by the World Health Organization

- 1. Purpose of this certificate:** To be introduced to Saudi Food and Drug Administration of Kingdom of Saudi Arabia for the purpose of Registration.
- 2. Certificate No.:** 1239/2021
- 3. Name and address of site:** Orchidia Pharmaceutical Industries (Al Obour City – north industrial zone extension, Parts No. (14,15)- Block no. (12011).  
• On the basis of the inspection, we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in the Table 1 below
- 4. Manufacturer's license number:** 112019041100002 (License issued from Industrial Development Authority)

Dosage form(s)	Last Inspection	Category(ies)	Activities carried out by the company
<u>Sterile:</u> Eye gel – Contact lenses solution	2019	General Category(ies)	Production and packaging
<u>Sterile:</u> Eye drops – Eye drops (single unit dose)- Eye drops (Solution – suspension – gel – emulsion)	2020		

- The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.
- This certificate remains valid until 9/12/2022 It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Dr. Amira Mahgoub  
*Amira Mahgoub*  
2021  
Manager of the Inspection Administration of  
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Pharmaceuticals Factories.



Authenticated  
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