



Serial :0009/2024

Licensing inspection report summary

Part 1: Manufacturer details:

- **Manufacturer name:** Super Care Pharmaceutical Industries
- Manufacturer address:** : Plots no .135 -industrial zone – Motobus- Kafr -Elsheikh
- **New manufacturer:** √ **- licensed manufacturer:** ×
- **Licensing inspection date:** 29/4/2024
- **Date of previously licensing inspections:**

Part 2: Scope of licensing inspection

Production line of Solid dosage forms includes powder filled in: Sachets/ glasses bottles with aluminum cover- (Plastic /Cans) in plastic sachet or not (Vet).

Part 3: Brief description about previously licensed production lines (in case of licensed factory)

- Super Care Pharmaceutical Industries is a new factory its production lines as described

Part 4: Summary of the findings and comments

- The opening meeting started with a presentation explaining the manufacturer activities in details from the authorized person who represented the factory layout including the different flows as personnel, material, waste, areas classification and differential pressure.
- Then a tour for the facility was conducted to involve production areas, as well as service areas, warehouses, and quality control laboratories.
- After the tour, the required documents for area and equipment qualification and all the required documents were reviewed.
- A close meeting was held by the committee members to decide the final conclusion and the committee decision was taken.
- Wrap up meeting was held to inform the factory representatives with the committee final decision

Part 5: Areas inspected

Preparation area, filling area, packaging areas, AHUs , laboratories ,warehouse , water treatment station, compressed air generator, all supplies



Part 6: Description

1- "Preparation area, filling area, packaging area, all supplies water treatment station , compressed air generator "

The facility shows compliance to GMP guidelines regarding:

- Suitable layout showing logic process flow.
- The factory premises shows adequate spaces .
- Classification and differential pressure were revised and complying.
- Suitable equipment used in manufacturing process.
- Facility was kept clean and had adequate lighting, ventilation, and environmental control.
- Area and equipment documentations and qualification were revised.
- Area supplies qualifications were revised and found complying.

2- Warehouse

- Adequate spaces and segregation of quarantine and released for raw materials, finished products , packaging materials was present.

Warehouses were found in a good cleanliness state with adequate lighting, ventilation.

3- Laboratories

- Laboratories premises layout was designed to suit the operations to be carried out.
- Appropriate calibrated equipment were present.
- Sufficient space was provided

Part7: References

- As per the law 151 for year 2019 of "promulgating law establishing the Egyptian authority for unified procurement, medical supply and technology management (AUPP) and Egyptian drug Authority (EDA) article 17 which stated "EDA shall exercise all regulatoryaccording to international standards".
- Also, as per prime minister degree no.777 for year 2020 article 17 which stated "...EDA adoption of standards and requirements of world health organization for the norms and requirements of good manufacturing practice (GMP).
- And all with taking into considerations the WHO references listed in the following link:
- <https://www.edaegypt.gov.eg/ar/%D8%A7%D9%84%D9%82%D9%88%D8%A7%D9%86%D9%8A%D9%86-%D9%88%D8%A7%D9%84%D9%82%D8%B1%D8%A7%D8%B1%D8%A7%D8%AA->



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Part 8 : Conclusion & The licensing inspection committee final decision.

Conclusion:

-Based on the areas inspected, the people met, and the documents reviewed, have been conducted at an acceptable level of compliance regarding : Production Areas , Equipment , warehouses , laboratories , Utilities , reviewed documents.

The licensing inspection committee final decision.

Granting the license . .