



PHARMA DEVILS

**OPERATIONAL QUALIFICATION PROTOCOL CUM
REPORT FOR
BIOMETRIC SYSTEM**

PROTOCOL No.:

**OPERATIONAL QUALIFICATION
PROTOCOL CUM REPORT
FOR
BIOMETRIC SYSTEM**

| | |
|--------------------------------|--------------------------------------|
| EQUIPMENT ID. No. | |
| LOCATION | MLT Area & Sterility Area |
| DATE OF QUALIFICATION | |
| SUPERSEDES PROTOCOL No. | Nil |



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**1.0 PRE – APPROVAL:
INITIATED BY:**

| DESIGNATION | NAME | SIGNATURE | DATE |
|--|-------------|------------------|-------------|
| OFFICER/EXECUTIVE (QUALITY ASSURANCE) | | | |

REVIEWED BY:

| DESIGNATION | NAME | SIGNATURE | DATE |
|--|-------------|------------------|-------------|
| OPERATING MANAGER (QUALITY ASSURANCE) | | | |
| HEAD (ENGINEERING) | | | |
| HEAD (QC-MICROBIOLOGY) | | | |

APPROVED BY:

| DESIGNATION | NAME | SIGNATURE | DATE |
|-------------------------------------|-------------|------------------|-------------|
| HEAD (QUALITY ASSURANCE) | | | |



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2.0 OBJECTIVE:

- To verify that the equipment operates in accordance with the design and user requirements as defined by set acceptance criteria and complies with relevant cGMP Requirements.
- To verify the Operational features of Biometric System and to ensure that it produces desired Quality & rated output according to manufactures specifications.
- To verify all the Operational features from user point of view of the Equipment, Cleaning Procedure, Startup & Shut down Procedure and Safety Features.

3.0 SCOPE:

- The scope of this Operational qualification protocol cum report is limited to qualification of Biometric System.
- This Protocol will define the methods and documentation used to perform OQ activity of Biometric System successful, completion of this Protocol will verify that Biometric System meet all acceptance criteria and ready for Daily use.



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4.0 RESPONSIBILITY:

The Validation Group, comprising of a representative from each of the following departments, shall be responsible for the overall compliance of this Protocol cum Report:

| DEPARTMENTS | RESPONSIBILITIES |
|--------------------------|---|
| Quality Assurance | <ul style="list-style-type: none">• Preparation and Compilation of the Operational Qualification Protocol cum Report.• Co-ordination with QC-Microbiology and Engineering to carryout Operational Qualification.• Monitoring of Operational Qualification Activity.• Post approval of Operational Qualification Protocol Cum Report. |
| QC-Microbiology | <ul style="list-style-type: none">• Review & Pre Approval of Protocol cum Report.• To Co-ordinate and support for Execution of Qualification study as per Protocol.• Post Approval of Qualification Protocol Cum Report. |
| Engineering | <ul style="list-style-type: none">• Review & Pre Approval of Protocol cum Report.• Co-ordination, Execution and technical support in Biometric System Operational Qualification Activity.• Responsible for Trouble Shooting (if occurs during execution).• Post Approval of Qualification Protocol Cum Report. |



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5.0 EQUIPMENT DETAILS:

| | | |
|---------------------------------|------------------|--|
| Equipment Name | Biometric System | |
| Equipment ID. | | |
| Manufacturer's Name | | |
| Modal | | |
| Location of Installation | | |

6.0 SYSTEM DESCRIPTION:

Make: ZKT_ECO

- This manual introduces the operation of user interfaces and menu functions of Pro Capture-X Access Control terminal.
- The pictures in this manual may not be exactly consistent with those of your product; the actual product's display shall prevail.
- Not all the devices have the function with ★, the real product prevails.

7.0 PRE – QUALIFICATION REQUIREMENTS:

7.1 Verification of Documents:

- DQ Protocol Cum Report
- IQ Protocol cum Report
- SOP For Operation and Handling of Biometric System in Microbiology Laboratory.



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8.0 CRITICAL VARIABLES TO BE MET:

8.1 Verification of documents:

The results of any tests should meet the limits and acceptance criteria specified in the test documents.
Any deviations or issues should be rectified and documented prior to OQ commencing.

| S.No. | Document Name | Document/SOP No. | Completed (Yes/No) | Checked By (Engineering) Sign/Date |
|-------|--|------------------|--------------------|------------------------------------|
| 1. | DQ Protocol Cum Report | | | |
| 2. | IQ Protocol Cum Report | | | |
| 3. | SOP For Operation And Handling Of Biometric System in Microbiology Laboratory. | | | |

Checked By
(QC- Micro.)
Sign/Date:

Verified By
(Quality Assurance)
Sign/Date:

Inference:

.....
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.....

Reviewed By
(Manager QA)
Sign/Date:



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8.2 Operational and Functional Checks:

Operate the Biometric System as per Manufacturer's Manual/SOP and Check for the following functions of the equipment. The Equipment should function as desired.

| S.No. | Name of component /Accessory | Area | Physical Condition | Working | Discrepancy Yes/No | Checked by (Sign/Date) |
|-------|------------------------------------|------|--------------------|---------|--------------------|------------------------|
| 1. | Electro-magnetic Lock | | | | | |
| 2. | | | | | | |
| 3. | Biometric Systems | | | | | |
| 4. | | | | | | |
| 5. | Display working Time / Date | | | | | |
| 6. | | | | | | |
| 7. | Light glowing Green/ Red | | | | | |
| 8. | | | | | | |
| 9. | Voice indicator | | | | | |
| 10. | | | | | | |
| 11. | Door Opening | | | | | |



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| S.No. | Name of component /Accessory | Area | Physical Condition | Working | Discrepancy Yes/No | Checked by (Sign/Date) |
|-------|------------------------------|------|--------------------|---------|--------------------|------------------------|
| 12. | | | | | | |
| 13. | Exit Push Button | | | | | |
| 14. | | | | | | |

Acceptance Criteria: Access control gives beep with green light for authorized person. Unauthorized entry should be rejected with beep with red light. Light should glow.

Checked By
(QC-Micro)
Sign/Date:

Verified By
(Quality Assurance)
Sign/Date:

Inference:

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Reviewed By
(Manager QA)
Sign/Date:



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8.3 Power Failure Verification:

| Item | Acceptance Criteria | Observation | Observed By (Engineering) Sign/Date |
|-----------------------------|--|-------------|---|
| Main Power shut down | Equipment stops in safe and secure condition | | |
| Main Power Restored | Equipment can be restarted with no problems or adverse conditions. | | |

Checked By
(QC-Micro)
Sign/Date:

Verified By
(Quality Assurance)
Sign/Date:

Inference:

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.....
.....

Reviewed By
(Manager QA)
Sign/Date:

9.0 REFERENCES:

The Principle Reference is the following:

- Validation Master Plan.
- Design Qualification.
- Installation Qualification.

10.0 DOCUMENTS TO BE ATTACHED:

- Any Other Relevant Documents

11.0 DEVIATION FROM PREDEFINED SPECIFICATION IF, ANY:

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12.0 CHANGE CONTROL, IF ANY:

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13.0 REVIEW (INCLUSIVE OF FOLLOW UP ACTION, IF ANY):

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14.0 CONCLUSION:

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15.0 RECOMMENDATION:

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16.0 ABBREVIATIONS:

| | | |
|------|---|--------------------------------------|
| WHO | : | World Health Organization |
| cGMP | : | Current Good Manufacturing Practices |
| QA | : | Quality Assurance |
| IQ | : | Installation Qualification |
| OQ | : | Operational Qualification |
| EQ | : | Equipment |
| BMT | : | Biometric System |
| SOP | : | Standard Operating Procedure |



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17.0 POST APPROVAL:

| DESIGNATION | NAME | SIGNATURE | DATE |
|--|-------------|------------------|-------------|
| OFFICER/EXECUTIVE (QUALITY ASSURANCE) | | | |

| DESIGNATION | NAME | SIGNATURE | DATE |
|--|-------------|------------------|-------------|
| OPERATING MANAGER (QUALITY ASSURANCE) | | | |
| HEAD (ENGINEERING) | | | |
| HEAD (QC-MICROBIOLOGY) | | | |

| DESIGNATION | NAME | SIGNATURE | DATE |
|-------------------------------------|-------------|------------------|-------------|
| HEAD (QUALITY ASSURANCE) | | | |