

# HEALTH SCIENCES AUTHORITY

REGULATORY GUIDANCE

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## GUIDANCE NOTES ON GOOD DISTRIBUTION PRACTICE

Draft Document



## Introduction

This guide is intended for those involved in the storage, transportation and distribution of Active Pharmaceutical Ingredients (API), Therapeutic Products (TP), Chinese Proprietary Medicines (CPM), and Cell, Tissue and Gene Therapy Products (CTGTP) with its active substances and starting materials, collectively referred to herein as *products*. This guide can also be applicable for Clinical Research Materials (CRM). This guide applies to all steps in the distribution/supply chain.

The objective of the guide is to ensure that the quality and integrity of the products are maintained throughout the distribution chain. The manufacturers, importers and distributors share important and distinctive roles and responsibilities to ensure that products are of the required quality for their intended use.

This guide aims to describe the critical and important controls appropriate for the storage and distribution of these products. Not all of the controls described will be relevant to every situation. It is recognized that some of the controls are not applicable to certain companies or environment. The controls should be adapted to meet individual company's needs where necessary. The relevance of any control should be determined taking into consideration the specific risks the company is facing such that the desired standards of quality are achieved.

Cold chain products are defined as products which are registered with the requirement of cold chain management. For cold chain products, both the requirements as stipulated in this guide and Annex 1 will be applicable.

## 1 PERSONNEL

- 1.1 There should be an adequate number of trained personnel involved in all stages of the distribution/supply chain. The number of personnel required will depend on the volume and scope of activities.
- 1.2 All personnel should be competent and possessing appropriate knowledge and experience, and where applicable, the relevant professional and technical qualifications for the tasks assigned to them.
- 1.3 All personnel should receive initial and continuing training in relation to Good Distribution Practice standards, operating procedures and safety issues, in accordance with a written training procedure. Special training should be provided for personnel dealing with special categories of products such as cytotoxic,

infectious or sensitizing products, products presenting special risks of abuse (including narcotic and psychotropic substances), CTGTP and cold chain products. Training records should be maintained.

## 2 PREMISES AND EQUIPMENT

- 2.1 Premises should be designed or adapted to ensure that the required storage conditions are maintained. Premises should also have sufficient security to prevent unauthorized access and misappropriation of the products.
- 2.2 Receiving and dispatch bays should protect products from the weather. The receiving area should be designed and equipped to allow cleaning of the containers of incoming materials, if necessary, before storage.
- 2.3 Storage areas should be of sufficient capacity to allow orderly and segregated storage of the various categories of products: those in quarantine and released, rejected, returned or recalled products. These designated storage areas should be clearly marked and the access to the products in quarantine and those that are rejected, returned or recalled should be restricted to authorized personnel. Any system (e.g. computerized and bar coding system) replacing the physical separation should give equivalent assurance in segregation and restriction in accessibility.
- 2.4 The storage areas should have adequate lighting and ventilation to enable all operations to be carried out accurately and safely. Premises should be carefully maintained, ensuring that repair and maintenance operations do not present any hazard to the quality of the products.
- 2.5 Premises should be dry, clean and free of accumulated waste and dust. A written cleaning procedure should be available indicating the frequency and methods to be used to clean the premises. Cleaning should be conducted so as not to present a source of contamination. Cleaning records should be maintained. For cytotoxic, infectious, sensitizing products or CTGTP, there should be appropriate procedures for the cleaning up of any spillage to ensure complete removal of any risk of contamination.
- 2.6 Products should be stored off the ground and suitably spaced to permit cleaning and inspection. Pallets should be well maintained and kept in a good state of cleanliness.

- 2.7 Storage conditions for products should be in compliance with the instructions on the product label. All equipment impacting on storage and distribution of products should be designed, located, maintained and cleaned to a standard which suits its intended purpose. The storage areas should be equipped with recorders or devices that will continuously monitor the storage conditions and record the relevant readings such as maximum and minimum temperature and humidity of the day. Appropriate actions on the premises, equipment and/or products should be taken when the storage conditions are not met and these actions taken should be recorded.
- 2.8 The recorders and devices for monitoring the storage conditions should be located in areas that are most likely to show fluctuations and/or the hottest and coldest locations where appropriate. These measuring equipment should be calibrated for the required operating range at defined intervals. Calibration of these measuring equipment should be traceable to national or international standard and such calibration records should be maintained.
- 2.9 Appropriate and suitable storage conditions should be provided for hazardous, sensitive and dangerous products such as combustible liquids and solids, pressurised gases, highly toxic substances, CTGTP and radioactive products.
- 2.10 Premises should be designed and equipped to prevent the entry of insects, rodents and other pests/animals. There should also be a pest control programme to identify and prevent pest infestation. Appropriate records should be maintained.
- 2.11 The presence of food, drink, smoking material or medicinal products for personal use should be prohibited in the storage areas.

### **3 STOCK HANDLING, STOCK CONTROL AND DELIVERIES**

- 3.1 Upon receipt, each incoming delivery should be checked for tampering and damage. Label description, type and quantity of the incoming products should also be physically verified against the relevant purchase order information. If necessary, any container or the entire delivery should be quarantined or set aside for further investigation. The type and nature of checks should be stated in a written procedure.

- 3.2 Products subject to specific storage requirements (e.g. narcotics, cold-chain products, CTGTP) should be immediately identified and stored in accordance with the written procedure. Incoming containers should be cleaned if necessary, before storage. Any activities performed on the incoming goods should not impact on the quality of the products.
- 3.3 Products in cartons/bulk packs should be adequately labeled with at least the product name, the batch number and the expiry date or retest date.
- 3.4 Products with broken seals, damaged packaging or suspected of possible tampering/contamination must not be sold or supplied.
- 3.5 Periodic stock reconciliation should be performed comparing the actual and recorded product quantity. All significant stock discrepancies should be investigated to check for inadvertent mix-ups and wrong issuance of stocks.
- 3.6 Products bearing an expiry date must not be received or supplied after their expiry date or so close to their expiry date that this date is likely to occur before the products are being used by the consumer.
- 3.7 A system should be in place to ensure that products due to expire first are sold and/or distributed first (Earliest-Expiry-First-Out, EEFO). Where no expiry dates exist for the products, FIFO (First-In-First-Out) should be applied. Deviations may, however, be permitted in exceptional cases where such deviation is appropriate and justified.
- 3.8 Deliveries should be made only to wholesale dealers or persons who are authorized to supply the products.
- 3.9 A written procedure on the delivery of the products to customers should be available.
- 3.10 Products should be transported in such a way that:
  - a) their identification is not lost;
  - b) they do not contaminate, and are not contaminated by, other products or materials;
  - c) adequate precautions are taken against spillage, breakage or theft;
  - d) they are secure and not subject to unacceptable degrees of heat, cold, light, moisture or other adverse influence, nor to be attacked by microorganisms or pests.

- 3.11 The vehicle/mode of transportation should not be used as a store for the products.

#### **4 DISPOSAL OF PRODUCTS**

- 4.1 Products intended for destruction should be appropriately identified, held separately, and handled in accordance with a written procedure.
- 4.2 Destruction of products should be carried out in accordance with the national legislative and regulatory requirements and with due consideration to protect the environment.
- 4.3 Records of all disposed products should be retained for a defined period.

#### **5 DOCUMENTATION**

- 5.1 The documentation system should include the specifications of products (applicable mainly to importers), procedures, instructions, protocols, contract, records, Certificate of Analysis and data, in paper or in electronic form. These documents should be made available for audit and upon request by the licensing authority.
- 5.2 Written procedures should be available to describe the different operations which may affect the quality of the products or of the distribution activities: personnel training, receipt and checking of incoming products, storage, deliveries, cleaning and maintenance of premises, pest control, monitoring and recording of the storage conditions, security of stocks on site, withdrawal of saleable stock, returned products, complaints, recalls, self-inspection, audit and assessment of contract acceptor, etc. The associated records of actions taken or conclusions reached should be maintained.
- 5.3 The title, nature and purpose of each document should be clearly stated. Documents should be uniquely identifiable and the effective date should be defined. The contents of documents should be clear and unambiguous. All documents should be approved, signed and dated by an appropriate authorized person(s) and should not be changed without the necessary authorization.
- 5.4 Documents should be reviewed regularly and kept up to date. When a document has been revised, a system should exist to prevent inadvertent use of the superseded version.

- 5.5 Records should be made or completed at the time each action is taken in such a way that all significant activities or events are traceable. Handwritten data entry should be made in a clear, legible and indelible way. Any alteration made to the entry should be signed and dated, and the alteration should permit the reading of the original information.
- 5.6 A record of receipt and distribution of the products shall be kept, stating the product name, date of transaction, invoice/delivery order number, name and address of purchaser/supplier, batch number, expiry date, quantity received/sold and stock balance. Relevant documents for the distribution (including records for samples) should be maintained.
- 5.7 Documents should be retained for a duration as in accordance with the legal requirements and be readily retrievable.
- 5.8 Each employee should have ready access to all necessary documentation for the tasks executed.
- 5.9 Data may be recorded by an electronic data processing system but detailed procedures relating to the system in use should be available and the accuracy of the records should be checked. Only authorized persons should be able to enter or modify data in the computer and there should be a record of changes and deletions (i.e. audit trail); access should be restricted by password or other means. It is particularly important that the data, including audit trail, are readily available throughout the period of retention. These data should also be protected by back-up transfer on separate hard disc, paper or other means.

## **6 PRODUCT COMPLAINTS**

- 6.1 There should be a written procedure describing the actions to be taken in the handling of all written and oral complaints regarding a possible product defect. There should be a record for each individual product complaint.
- 6.2 The procedure for handling product complaints shall ensure that the complaints received are investigated and followed through, and that corrective actions are taken to prevent recurring problems. All original details of the product complaint, investigations and subsequent corrective and preventive actions taken, including

product recall should be documented in the product complaint record.

- 6.3 Within the company, a person shall be designated to handle product complaints. This person must have the authority to initiate investigations.
- 6.4 If a product defect is discovered or suspected in a batch, consideration should be given to determine whether other batches are also affected.
- 6.5 Product complaint records should be reviewed regularly for any indication of specific or recurring problems requiring attention.

## **7 PRODUCT RECALL**

- 7.1 An emergency plan for urgent recalls and a non-urgent product recalls procedure should be described in writing.
- 7.2 A person or committee should be designated for the co-ordination and execution of all product recalls.
- 7.3 In the event of a product recall, all customers to whom the product has been distributed shall be informed with the appropriate degree of urgency. The recall message should indicate whether the recall should be carried out at the retail level, and whether there is a need to remove all recalled products immediately from the shelves and prevent their mixing with other saleable stocks.
- 7.4 The local regulatory authority should be informed of all product recalls. If the product is exported, the overseas counterparts and/or regulatory authorities must be informed of the recall.
- 7.5 Where product recall affects a particular batch, consideration should also be given to determine whether other batches are also affected.
- 7.6 All actions taken in connection with the product recall must be approved by the company and/or regulatory authorities and recorded.
- 7.7 The progress of recall process should be recorded and a final report issued, which includes reconciliation made between delivered and recovered quantities of products.



## 8 RETURNED PRODUCTS

- 8.1 There should be a written procedure describing the handling of returned products and the corresponding records of all returned products should be kept.
- 8.2 All returned products should be kept apart from saleable stock to prevent redistribution until a decision has been reached regarding their disposition.
- 8.3 Returned products should only be returned to saleable stock if all of the following are confirmed:
- a) the products are in their original unopened and undamaged secondary packaging and are in good condition;
  - b) it is known that the products have been transported, stored and handled under proper conditions;
  - c) the remaining shelf life period is acceptable; and
  - d) the products have been examined and assessed by appropriate and qualified personnel. This assessment should take into account the nature of the product, any special storage conditions required, and the time which had elapsed since it was distributed. Special attention should be given to thermolabile products. Advice should be sought from the marketing authorization (product licence) holder or manufacturer as necessary.

Where any doubt arises over the quality of the product, it should not be considered suitable to be returned to saleable stock.

- 8.4 The returned products should be formally released to saleable stock by a nominated, responsible person following a satisfactory quality re-evaluation.
- 8.5 Products returned to saleable stock should be placed in accordance with the FEFO or FIFO system.

## 9 COUNTERFEIT PRODUCTS

- 9.1 The sale and distribution of a suspected counterfeit product should be suspended immediately.
- 9.2 Any counterfeit products found in the supply chain should be physically segregated from other materials to avoid any confusion. They should be clearly labeled as "Not for Sale" or with other

similar phrases/words. All relevant activities in relation to such products should be documented and records retained.

- 9.3 The regulatory authority and the holder of the marketing authorisation of the original product should be informed immediately.
- 9.4 Upon confirmation as a counterfeit product, a formal decision should be taken on removal of such product from the market, ensuring that it does not re-enter the supply chain, including retention of any samples necessary for public health, regulatory, or legal needs and arrangements for its disposal. All related decisions should be appropriately documented.

## **10 SELF-INSPECTION**

- 10.1 Self inspections should be conducted to monitor the implementation and compliance with this (GDP) standard and to propose necessary corrective and preventive measures.
- 10.2 Self-inspections should be conducted in an impartial and detailed way by designated, competent personnel. There should be a written procedure on self-inspection stating the persons involved in self-inspection, the frequency of self-inspection and the inspection criteria.
- 10.3 All self-inspections should be recorded. This record should include all observations made during the inspection. In the event that irregularities and/or deficiencies are observed, their cause should be determined and the corrective and preventive actions (CAPA) should be documented and followed up.

## **11 CONTRACT ACTIVITIES**

- 11.1 Any activity covered by the GDP Guide that is outsourced should be defined, agreed and controlled in order to avoid ambiguity, which could affect the quality of the product. There must be a written contract between the contract giver and the contract acceptor which clearly establishes the roles and responsibilities of each party, including compliance with this standard.
- 11.2 The contract giver is responsible for assessing the competence of the contract acceptor to successfully carry out the work required and for ensuring by means of the contract and through audits that

the principles and guidelines of GDP are followed. An assessment of the contract acceptor should be performed before commencement of, and whenever there has been a change to, the outsourced activities. Any contract acceptor should be audited periodically by the contract giver. The requirement for audit and the frequency for audit should be defined based on risk depending on the nature of the outsourced activities. Audits by the contract giver should be permitted at any time.

- 11.3 The contract giver should provide the contract acceptor with all the information necessary to carry out the outsourced operations in accordance with the specific product requirements and any other relevant requirements.
- 11.4 The contract acceptor should have adequate premises and equipment, procedures, knowledge and experience, and competent personnel to carry out the work ordered by the contract giver.
- 11.5 The contract acceptor should not pass to a third party any of the work entrusted to him under the contract without the contract giver's prior evaluation and approval of the arrangements. Such arrangements should ensure that the wholesale distribution information is made available in the same way as between the original contract giver and contract acceptor.
- 11.6 The contract acceptor should refrain from any activity which may adversely affect the quality of the product(s) handled for the contract giver.
- 11.7 The contract acceptor must forward any information that can influence the quality of the product(s) to the contract giver in accordance with the requirement of the contract.

## **12 HANDLING OF CELL, TISSUE AND GENE THERAPY PRODUCTS & RELATED SUBSTANCES / MATERIALS**

- 12.1 There must be an effective, current risk-based quality system which adequately assesses the risks that impacts product quality; considers the proper controls and mitigation measures; as well as putting in place the appropriate communications and active process reviews, to ensure maintenance of appropriate storage and distribution conditions for the CTGTP and its related substances and materials (hereinafter known as "products"). The level of effort, formality and documentation should commensurate with the level of risk and fit for its purpose to the protection of the patient.

- 12.2 The appropriate use of risk-based processes and documentation should demonstrate the company's obligation and compliance with regulatory requirements, through appropriate communications of these documented evidences between the company and the regulatory authority.
- 12.3 The established infrastructure should build in relevant capability(ies) that permit real-time monitoring of the conditions, appropriate alert mechanisms, and accurate track-and-trace of the CTGTP (including that of the cells/tissues) through storage, transport and delivery in the supply chain. All these tasks should be carried out by trained personnel.
- 12.4 The data generated from the system should be reviewed and maintained.
- 12.5 The system should ensure that traceability data may be accessed rapidly in case of an adverse reaction from the patient.
- 12.6 The data are to be retained for a minimum of 30 years after the expiry date of the product, or a longer period if provided for in the product registration.
- 12.7 Storage containers should be sealed, clearly labelled and kept at an appropriate temperature. When liquid nitrogen is used, liquid-nitrogen vapour phase storage may be associated with vertical temperature gradients. Temperature gradients should be monitored and vertical temperature gradients should be minimised. The liquid nitrogen level should be monitored. Deviation from set limits and corrective and preventive action taken should be recorded. A stock inventory and the relevant monitoring records must be kept.
- 12.8 There should be written procedures established to ensure that adequate conditions have been maintained before products that are removed from the storage area (e.g. cryostorage) can be returned to storage area. All relevant activities in relation to such products should be documented and records retained.
- 12.9 There should be written procedures and documentation in place for the secured handling and storage of products with positive serological markers (e.g. Hepatitis B, HIV, etc.) or viral vector products. This should describe the storage arrangements, such as the type of storage chamber, placement and retrieval process, as well as the cleaning processes to minimise the risk of cross-

contamination, while maintaining the quality of the products and facilitate accurate product retrieval.

- 12.10 Particular attention should be paid to implementing appropriate measures and labelling to prevent mix-ups of products (e.g. autologous products, products intended for specific patients).
- 12.11 There should be an emergency plan for dealing with accidental spillage of viable organisms. This should consider the methods and procedures for containment, protection of personnel, cleaning, decontamination, waste management and safe return to use.
- 12.12 CTGTP should be transported in accordance with the conditions defined in the product registration or clinical trial authorisation (where applicable).
- 12.13 The conditions during distribution and delivery of CTGTP (e.g. transportation routes, temperature, time limit, type of container) should be clearly defined and validated. All containers and packages need to be validated and suitable for the intended purpose.

### **13 HANDLING OF ACTIVE PHARMACEUTICAL INGREDIENT OR INTERMEDIATES**

- 13.1 This section are additional requirements which are relevant the agents, brokers, traders or distributors, generally referred to as “dealer” who may trade and/or take possession, distribute or store an API or intermediate.
- 13.2 Active Pharmaceutical Ingredient (API) is any substance or mixture of substances intended to be used in the manufacture of a drug (therapeutic) product and that, when used in the production of a drug, becomes an active ingredient of the drug product. Such substances are intended to furnish pharmacological activity or other direct effect in the diagnosis, cure, mitigation, treatment, or prevention of disease or to affect the structure and function of the body.
- 13.3 Intermediate is a material produced during steps of the processing of an API that undergoes further molecular change or purification before it becomes an API.
- 13.4 The dealers should maintain complete traceability of APIs and intermediates that they distribute. Documents that provides

traceability includes identity and address of the original manufacturer, purchase orders, transportation documentation, manufacturer's batch number, transportation and distribution records as well as authentic Certificates of Analysis.

- 13.5 Original certificates of analysis issued by the manufacturer or authenticated copies of the original certificates of analysis should be provided for each batch of intermediates or APIs on request by the customers.

## 14 REFERENCES

1. World Health Organization (WHO) Good Distribution Practices for Pharmaceutical Products. WHO Technical Report Series, No. 957, 2010.
2. Pharmaceutical Inspection Co-operation Scheme (PIC/S) Guide to Good Distribution Practice (GDP) for Medicinal Products.
3. HSA Guidelines on Good Manufacturing Practice for Cell, Tissue and Gene Therapy Products
4. USP <1044> Cryopreservation of Cells

## Annex 1

### COLD CHAIN PRODUCTS

1. There should be written procedures established to ensure that incoming cold chain products are delivered under the storage conditions in compliance with the instructions on the product label, which are based on the results of stability testing. Companies may make use of temperature data loggers or other temperature recording instruments to verify that the desired temperature has been maintained during delivery for each consignment received. Alternatively, there should be a simulation study carried out to validate the delivery conditions, taking into consideration the possible worst-case situation. This aforementioned requirement should also be applicable to the outbound deliveries.
2. The list of cold chain products should be made available for reference by staff handling the receipt of such goods and other relevant store personnel.
3. Cold chain products should be immediately identified upon receipt and stored under the storage conditions in compliance with the instructions on the product label. This should be carried out in accordance with the written procedure.
4. The subsequent checking for signs of tampering, damage and non-compliance with cold chain storage condition, as well as physically verifying the label description, and product quantity, against the relevant information in the purchase order should be carried out under the storage conditions as recommended on the product label unless otherwise justified.
5. All cold chain products (e.g. released, quarantined) must be stored under the storage conditions as indicated on the product label other than products to be disposed.
6. The cold room, which is used for storage of cold chain products, should be subject to temperature mapping studies, under representative conditions in order to identify the suitable locations for placing the temperature probes. The mapping exercise should be repeated accordingly if there are significant changes.

7. The temperature conditions of the cold room or refrigerator should be monitored and recorded on a continuous basis. The temperature probes should also be subject to periodic calibration for the required operating range. A regular maintenance programme should be established and carried out for the air conditioning system of the cold room and refrigerator. For storage units that has an auto-defrost facility, precautionary steps should be taken to ensure that the storage temperature is not affected during the defrost cycles.
8. The cold room or refrigerator should be installed with an alarm system to alert the staff to any temperature excursions. Action and alert limits should be established. The function of the alarm system should be subject to periodic testing.
9. Backup power should be available for the cold room to ensure that storage temperature conditions will be maintained and temperature probes and recording devices will continue to function in the event of a power failure. Any backup generators used should be subject to periodic testing. Alternative back-up plans that provide equivalent storage temperature conditions and monitoring system can be considered in the absence of backup generator.
10. There should be written procedures to describe the packing materials required, the packing configuration of transportation containers of cold chain products and labeling requirements that easily identify these products as products that require special delivery/storage conditions. The packing operation for the cold chain products should be recorded and there should be an independent check to ensure that the packing operation is carried out in accordance with the written procedure. The individual who carried out this independent check should initial in the packing records.
11. Refrigerated vehicles or transportation containers for cold chain products should be mapped and monitored if they provide the primary means for environmental control. However, this is not necessary if a qualified/validated insulated container is used for this purpose. Special care should be exercised when using dry ice during transportation. Products should not come into contact with dry ice as this may cause freezing of the product.
12. There should be written procedures available for the delivery of cold chain products. The planning of the delivery route should be taken into consideration.
13. There should be procedures established for handling temperature excursions that may occur during receiving, storage and delivery.



14. There should be procedures established to describe how product return requests should be handled and the disposition of such products.
15. Written contract(s) should be established to describe arrangements and responsibilities between the contract giver and contract acceptor with respect to contract warehousing and transportation of cold chain products. The contract should provide all the necessary information and define the conditions required for the contracted operations to be carried out, as well as the responsibilities of each party.
16. There should be written procedures available and appropriate training provided for all staff involved in the handling, receipt, storage, packing and delivery operations that may affect the quality of cold chain products.

**Annex 2**

General points to consider for auditor and auditee

<ul style="list-style-type: none"> <li>• <b><u>General Information</u></b> <ul style="list-style-type: none"> <li>- Any contract warehouse(s)</li> <li>- Approval available for the use of the warehouse</li> </ul> </li> <li>• <b><u>Personnel</u></b> <ul style="list-style-type: none"> <li>- Name and designation of personnel</li> <li>- Training programme and records</li> </ul> </li> <li>• <b><u>Premises and Equipment</u></b> <ul style="list-style-type: none"> <li>- Floor area</li> <li>- Layout plan</li> <li>- Store approval</li> <li>- Prevent unauthorized access</li> <li>- Adequate storage area with segregations</li> <li>- Appropriate for the products</li> <li>- Lights/ventilation</li> <li>- Dry and clean</li> <li>- Cleaning procedure</li> <li>- Cleaning records</li> <li>- Storage off ground</li> <li>- Storage – Sunlight</li> <li>- Thermometer / hygrometer and records</li> <li>- Warehouse design prevents pest entry</li> <li>- Appropriate pest control programme</li> </ul> </li> <li>• <b><u>Stock Handling, Stock Control and Deliveries</u></b> <ul style="list-style-type: none"> <li>- Receiving procedure</li> <li>- Appropriate types of checks conducted</li> <li>- Goods are labelled</li> <li>- EEFO / FIFO</li> <li>- Stock Reconciliation</li> <li>- Procedure for delivery of products</li> </ul> </li> <li>• <b><u>Disposal</u></b> <ul style="list-style-type: none"> <li>- Written procedure and records</li> </ul> </li> <li>• <b><u>Documentation</u></b> <ul style="list-style-type: none"> <li>- Procedure for stock handling and labeling and monitoring of storage conditions</li> <li>- SOP signed and formalized</li> <li>- Content of SOP clear and kept up to date</li> <li>- System to prevent inadvertent use of obsolete procedures</li> <li>- Receiving and distribution records, invoices and delivery orders</li> <li>- Record retention</li> <li>- Computerized record – restricted access, audit trail and back-up</li> </ul> </li> <li>• <b><u>Product Complaints</u></b> <ul style="list-style-type: none"> <li>- SOP and records</li> <li>- System for investigation and review</li> </ul> </li> <li>• <b><u>Product Recall</u></b> <ul style="list-style-type: none"> <li>- SOP and records</li> <li>- Designated person</li> <li>- Level of recall established</li> </ul> </li> <li>• <b><u>Returned Products</u></b> <ul style="list-style-type: none"> <li>- SOP and records</li> <li>- Assessment criteria</li> <li>- Authorization for re-sale</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• <b><u>Self-Inspection</u></b> <ul style="list-style-type: none"> <li>- SOP and records</li> </ul> </li> <li>• <b><u>Contract Activities</u></b> <ul style="list-style-type: none"> <li>- Contract</li> <li>- Content define responsibilities and requirements</li> </ul> </li> <li>• <b><u>Poisons &amp; CD Requirements</u></b> <ul style="list-style-type: none"> <li>- Sample control record</li> <li>- Signed orders/invoices/other supporting documents</li> <li>- CD register</li> </ul> </li> <li>• <b><u>Cytotoxic Products</u></b> <ul style="list-style-type: none"> <li>- Labels for identification and warning</li> <li>- Appropriate training</li> <li>- Procedure for dealing with spillage incident</li> <li>- Cytotoxic spillage control kit</li> </ul> </li> <li>• <b><u>Cell, Tissue and Gene Therapy Products</u></b> <ul style="list-style-type: none"> <li>- Quality risk management</li> <li>- Real-time monitoring of storage, transport and delivery conditions</li> <li>- Track-and-trace of CTGTP</li> <li>- Rapid access to traceability data</li> <li>- Data retention period</li> <li>- Handling of storage containers</li> <li>- Procedures for return of CTGTP back to storage</li> <li>- Procedures for handling of CTGTP with positive serological markers or viral vector products</li> <li>- Prevention of mix-ups</li> <li>- Emergency plan for dealing with spillage</li> <li>- Defined transportation conditions</li> <li>- Validated containers and packages used for delivery</li> </ul> </li> <li>• <b><u>Active Pharmaceutical Ingredients</u></b> <ul style="list-style-type: none"> <li>- Traceability</li> <li>- Certificate of Analysis</li> </ul> </li> <li>• <b><u>Cold Chain Products</u></b> <ul style="list-style-type: none"> <li>- Receiving and incoming checks</li> <li>- Labels/means to identify cold chain products</li> <li>- Temperature mapping</li> <li>- Thermometer and records</li> <li>- Maintenance programme</li> <li>- Alarm system for temperature excursion</li> <li>- Backup generator/plan</li> <li>- Packing procedure and records, and independent check</li> <li>- Temperature mapping for vehicles or qualified/validated containers</li> <li>- Monitoring of storage conditions during transportation or simulation study</li> <li>- Delivery procedure</li> <li>- Procedure to handle temperature excursion</li> <li>- Procedure for handling returned products</li> <li>- Contracts</li> <li>- Training programme and records</li> </ul> </li> </ul>
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**END OF DOCUMENT**

# HEALTH SCIENCES AUTHORITY

Health Products Regulation Group  
Blood Services Group  
Applied Sciences Group

[www.hsa.gov.sg](http://www.hsa.gov.sg)

**Contact:**

GDP Unit  
Audits Branch  
Audit & Licensing Division  
Health Products Regulation Group  
Health Sciences Authority

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