



## **BAGADIONG, FERMIN JOEL JR. SEGOVIA**

Contact Details: *Mobile no.: (+65) 8511 2869 | email: bagoJOONg@yahoo.com*

***A Registered Pharmacist in the Philippines; Certified ISO 13485 and ISO 9001 Internal Auditor with working experience in healthcare services both in government (Regulatory, Hospital Pharmacy and Public Health) and private sector (Quality Assurance and Retail Pharmacy) and has worked in a GxP (GMP/GDP) certified facility.***

***Trained in ISO9001:2015, ISO13485:2016, GMP and GDP (SS620:2016) with an excellent track record of work performance, achieved all internal and external audits; initiated and developed training program.***

### **PROFESSIONAL QUALIFICATIONS:**

- ❖ **ISO 13485:2016 Internal Auditor**, 05 Apr 2019 (SGS International Certification Services Singapore Pte Ltd)
- ❖ **ISO 9001:2008 Internal Auditor**, 23-24 Jun 2014 (DHL Country QA)
- ❖ **ISO 13485:2003 Internal Auditor**, 20-22 Mar 2013 (SGS International Certification Services Singapore Pte Ltd)
- ❖ **Certified Pharmacy Technician Programme**, 27 Oct 2009 – 16 Nov 2009 (Pharmacy Technician Programme Guardian Health & Beauty (Singapore))
- ❖ **Registered Pharmacist**, July 1995 (Pharmacist Licensure Examination Professional Regulation Commission (PRC, Philippines))

### **EMPLOYMENT HISTORY:**

#### **1) SENIOR QUALITY ASSURANCE OFFICER**

##### ***DCH Auriga Singapore***

24 Penjuru Road, #03-04/#04-04 ALOG Commodity Hub, Singapore 609128

*Inclusive Date: 21 Dec 2020 to Present*

##### Job Description/Responsibilities:

- ❑ Document Controller, Compliance and Audit
  - Document Controller for DCH Quality Management System to ensure current work processes are compliance with ISO 9001, ISO 13485 standards and GMP/GDP requirements
  - Responsible for the maintenance and filing of Quality related reports and records
  - Engage with process owners using CAPA process to eliminate nonconformities and prevent recurrence in the quality system
  - Escalate and report to Management where systemic failures or regulatory non-compliance in the quality management system that are at risk
  - Ensure environmental controls, temperature monitoring, pest control and cleaning complies with requirements; Review of Calibration Reports; Review of Pest Control reports monthly.
  - Responsible for managing Annual Customer Survey.
  - Manage FMEA and Risk Assessment activities.
  - Conduct periodic SOP / WI training related to document control, redressing, QA or FMEA to ensure new hires are equipped with the control of the processes
  - Conduct refresher training when necessary or when there are changes in SOP/WI related to QA purview.
  - Conducts weekly Briefing (Quality sharing)
  - Assist in the suppliers' evaluation before suppliers are qualified in the approved vendor list
  - Manage complaints, investigation process as well as evaluate and present annual customer feedback survey.
  - Participate in continuous improvement activities, self-inspection program to drive compliance and efficiency improvement
  - Management Representative of the company
  - Lead Auditor for internal audits.
  - Key personnel for external and principals' audits
  - Support RFQ – review of QTA



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- ❑ Secondary Assembly
  - Designated Quality personnel in the GMP Secondary Assembly Operations.
  - Responsible for quality oversight of the Secondary Assembly Operations.
  - Perform inspection, sampling of products, batch records checking and final release of repackaged products and incoming printed packaging materials for secondary assembly operations.
  - Coordinate secondary assembly activities
  - Responsible for the ordering of printed packaging materials such as labels and cartons, needed for Secondary Assembly Operations.
  - Responsible for the stocks, issuance and reconciliations of the labels in DC warehouse.
  - Draft/create/revise Master Batch Packaging Record (MBPR)
  - Prepare and submit monthly KPI report
  - Supervise QC (Redressing Team Lead)
  - As Back-up QC
- ❑ Employee Engagement and Leadership
  - Provide coaching and leadership to the team.
  - Implement work improvement plan or introduce disciplinary action when situations required.
  - Conduct direct report annual performance appraisal and mid-year review

## **2) QUALITY EXECUTIVE**

***DB Schenker Singapore (Pte) Ltd*** (Department: **CL/SCM-Becton Dickinson RDC**)

5 Tuas View Lane, Singapore 637291

*Inclusive Date: 09 Sep 2019 to 18 Dec 2020*

### Job Description/Responsibilities:

- Program/department Site Quality
- Implement and monitor site's QMS in compliance with ISO (ISO 9001 and ISO 13485),
- Conduct checks on documents and processes to ensure procedures are followed
- Work with OPS Executive to resolve quality issues
- Prepare, review and implement change requests (CR) raised
- Create, revise, review Work Instructions
- Review and approve Quality related documentation
- Closure of NCRs / CAPAs (8D) / customer complaints / audit actions
- Training of staff in GDPMDS
- Conduct Weekly Briefing
- Review temperature related document
- Key personnel for internal and external audits
- Liaise with customer on quality related matter
- Attend Management Review (MR) and customer meetings
- Verify line clearance for labeling
- Verify contents and quality of print of labels
- Perform sampling check at labeling process
- Review and approve the labeling work orders
- Release of labeled products for packing
- Approve or reject the final batch secondary assembly and over-labeling product
- Supervise QC
- As back-up QC



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### **2) QUALITY ASSURANCE EXECUTIVE (QA Lead)**

**DHL Supply Chain Singapore Pte Ltd** (Department: **MEDTRONIC**)

Advanced Regional Centre, Tampines LogisPark, 1 Greenwich Drive, ARC Warehouse  
Block 1 Level 2, Singapore 533865

*Inclusive Date: 11 Sep 2017 to 04 Sep 2019*

Job Description/Responsibilities: *Carry out quality assurance and control related activities to ensure the business is operating to the highest standards of regulatory and quality compliance in the Life Sciences and Healthcare sector operations to guarantee patient safety.*

#### *A. Policies and Procedures*

- Ensure that Company Policies and Quality Procedures are effectively carried out.

#### *B. Quality Management System*

- Implement and monitor site's QMS in compliance with ISO (ISO 9001 and ISO 13485), Regulatory and GDPMDS requirements.

#### *C. Documentation Management*

- Ensure all QMS related documents are current and up to date.
- Follow-up with process owner to update documents timely.
- Update site QA Work Plan holistically and escalate to Quality Manager if there is gap in the plan versus completion of activity.
- Create, revise, review and approves SOP/work instructions.

#### *D. Audit Support*

- Facilitate internal and external audits at site level.
- Support/conduct intra-company quality audits directed by Country QA Lead.

#### *E. Self-Inspection*

- Performs self-inspection to ensure GxP and relevant certifications compliance.

#### *F. Risk Management*

- Lead and provide guidance on risk management team discussions.
- Perform risk assessments using FMEA tool

#### *G. Change Control*

- Ensure Change Request is raised for any changes that may have impact onto the quality of products
- Perform Change Impact Analysis

#### *H. Training*

- Site GxP trainer.
- Site induction program.
- Develop training materials.
- Conducts weekly briefing.
- Control training matrix and records.
- Close competency gaps by facilitating site quality related training.

#### *I. Facility Environmental Monitoring*

- Ensure environmental controls, temperature monitoring, pest control and cleaning complies with requirements. To raise deviations if needed.
- Maintains Masterlist of Equipment and ensure all equipment is calibrated on time.
- Conducts GEMBA walk for 5S Housekeeping and Safety.
- Review and verifies deviation reports.

#### *J. Corrective Action Preventive Action (CAPA)*

- Conduct meetings with CAPA owners/customers to drive closure
- Perform root cause analysis (RCA) with CAPA owners to determine the root cause
- Maintains CAPA Log

#### *K. Supplier Qualification*

- Conduct Desktop Audit
- Conduct on-site audit

#### *L. Site Quality Practitioner*

- Conduct annual quality self-assessment using the Path to Quality (PtQ) Effectiveness Assessment Checklist/MQS standards



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- Conduct pre-GoLive assessment
  - Gathers objective evidence via assessment interviews and maintains an audit trail.
  - Present assessment report and findings during Monthly Management Review.
  - Drive and ensure closure of all corrective and improvement action items, as a result of non-conformities raised following the assessment.
  - Support the Regional/Country Quality Team in fulfilling the objectives of the QA function.
  - Maintains, review and updates Site Manual.
  - Manage and maintain CAPA report system includes trending and reporting
- M. *QA/QC activities relating to production*
- Perform In-Process and Final Check of labeled products via AQL sampling.
  - Supervise Labeling Operators in labeling operation.
- N. *Support to Operations*
- Lead and provide guidance to the operations team on quality related matters.
  - Review Non-Conformance Reports and sign off for QA disposition.
  - Attends Daily OPS Meeting with the customer.
  - Back-up QC for labeling operations
  - Ensure RCA is performed to determine the root cause in NCR/CAPA responses
  - Ensure CAPAs are responded/closed on time
  - Control access (Key) to Quarantine Cage and Incubators/Fridge
  - Drive and ensure monthly Management Review is conducted.
  - Arranges and facilitates monthly disposal
  - Verifies on-hold products (PHO)
  - Respond to customers quality related issues in the warehouse
  - Perform other functions assigned by LSHC QA Manager and/or LSHC Director/General Manager

### **3) QUALITY ASSURANCE EXECUTIVE**

***DHL Supply Chain Singapore Pte Ltd*** (Department: **GE Healthcare**)

Advanced Regional Centre, Tampines LogisPark, 1 Greenwich Drive, ARC Warehouse  
Block 2 Level 2, Singapore 533865

*Inclusive Date: 13 Mar 2017 to 31 Jan 2018*

Job Description/Responsibilities: *Carry out quality assurance and control related activities to ensure the business is operating to the highest standards of regulatory and quality compliance in the Life Sciences and Healthcare sector operations to guarantee patient safety.*

#### **A. Policies and Procedures**

- Ensure that Company Policies and Quality Procedures are effectively carried out.

#### **B. Quality Management System**

- Implement and monitor site's QMS in compliance with ISO (ISO 9001 and ISO 13485), Regulatory and GMP requirements.

#### **C. Documentation Management**

- Ensure all QMS related documents are current and up to date.
- Follow-up with process owner to update documents timely.
- Update site QA Work Plan holistically and escalate to Quality Manager if there is gap in the plan versus completion of activity.
- Writes, revise, review and approves SOP/work instructions.

#### **D. Audit Support**

- Facilitate internal and external audits at site level.
- Support/conduct intra-company quality audits directed by Country QA Lead.

#### **E. Self-Inspection**

- Performs self-inspection to ensure GxP and relevant certifications compliance.

#### **F. Risk Management**

- Lead and provide guidance on risk management team discussions.
- Perform risk assessments

#### **G. Change Control**

- Ensure Change Request is raised for any changes that may have impact onto the quality of products



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- Perform Change Impact Analysis
- H. *Training*
  - Site GxP trainer.
  - Site induction program.
  - Develop training materials.
  - Conducts weekly briefing.
  - Control training matrix and records.
  - Close competency gaps by facilitating site quality related training.
- I. *Facility Environmental Monitoring*
  - Ensure environmental controls, temperature monitoring, pest control and cleaning complies with requirements. To raise deviations if needed.
  - Maintains Masterlist of Equipment and ensure all equipment is calibrated on time.
  - Conducts GEMBA walk for 5S Housekeeping and Safety.
  - Review and verifies deviation reports.
- J. *Corrective Action Preventive Action (CAPA)*
  - Conduct meetings with CAPA owners/customers to drive closure
  - Perform root cause analysis (RCA) with CAPA owners to determine the root cause
  - Maintains CAPA Log
- K. *Supplier Qualification*
  - Conduct Desktop Audit
  - Conduct on-site audit
- L. *Site Quality Practitioner (QP – Path to Quality)*
  - Conduct monthly self-assessment using the Path to Quality (PtQ) Effectiveness Assessment Checklist.
  - Conduct self-inspection at least one process a month.
  - Conduct pre-GoLive assessment
  - Gathers objective evidence via assessment interviews and maintains an audit trail.
  - Present assessment report and findings during Monthly Management Review.
  - Drive and ensure closure of all corrective and improvement action items, as a result of non-conformities raised following the assessment.
  - Support the Regional/Country Quality Team in fulfilling the objectives of the QA function.
  - Maintains, review and updates Site Manual.
  - Manage and maintain CAPA report system includes trending and reporting
- M. *Support to Operations*
  - Lead and provide guidance to the operations team on quality related matters.
  - Attends OPS Meeting with the customer.
  - Ensure RCA is performed to determine the root cause in CAPA responses
  - Ensure CAPAs are responded/closed on time
  - Drive and ensure monthly Management Review is conducted.
  - Verifies products during monthly disposal
  - Respond to customers quality related issues in the warehouse
  - Perform other functions assigned by LSHC QA Manager and/or LSHC Director

#### **4) QUALITY ASSURANCE EXECUTIVE**

**DHL Supply Chain Singapore Pte Ltd** (Department: **Healthcare Hub**)

7 Changi South Street 2, #01-00 Xilin Districentre Singapore 486415

*Inclusive Date: 01 May 2013 to 12 Mar 2017*

**\*\*Promoted**

Job Description/Responsibilities: *Carry out quality assurance and control related activities to ensure the business is operating to the highest standards of regulatory and quality compliance in the Life Sciences and Healthcare sector operations to guarantee patient safety.*

##### **A. Policies and Procedures**

- Ensure that Company Policies and Quality Procedures are effectively carried out.



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- B. Quality Management System**
  - Implement and monitor site's QMS in compliance with ISO (ISO 9001 and ISO 13485), Regulatory and GMP requirements.
- C. Documentation Management**
  - Ensure all QMS related documents are current and up to date.
  - Follow-up with process owner to update documents timely.
  - Update site QA Work Plan holistically and escalate to Quality Manager if there is gap in the plan versus completion of activity.
  - Writes, revise, review and approves SOP/work instructions.
- D. Audit Support**
  - Facilitate internal and external audits at site level.
  - Support/conduct intra-company quality audits directed by Country QA Lead.
- E. Self-Inspection**
  - Performs self-inspection to ensure GxP and relevant certifications compliance.
- F. Risk Management**
  - Lead and provide guidance on risk management team discussions.
  - Perform risk assessments
- G. Change Control**
  - Ensure Change Request is raised for any changes that may have impact onto the quality of products
  - Perform Change Impact Analysis
- H. Training**
  - Site GxP trainer.
  - Site induction program.
  - Develop training materials.
  - Conducts weekly briefing.
  - Control training matrix and records.
  - Close competency gaps by facilitating site quality related training.
- I. Facility Environmental Monitoring**
  - Ensure environmental controls, temperature monitoring, pest control and cleaning complies with requirements. To raise deviations if needed.
  - Maintains Masterlist of Equipment and ensure all equipment is calibrated on time.
  - Conducts GEMBA walk for 5S Housekeeping and Safety.
  - Review and verifies deviation reports.
- J. Supplier Qualification**
  - Conduct Desktop Audit
  - Conduct on-site audit
- K. Site Quality Practitioner (QP – Path to Quality)**
  - Conduct monthly self-assessment using the Path to Quality (PtQ) Effectiveness Assessment Checklist.
  - Conduct self-inspection at least one process a month.
  - Conduct pre-GoLive assessment
  - Gathers objective evidence via assessment interviews and maintains an audit trail.
  - Present assessment report and findings during Monthly Management Review.
  - Drive and ensure closure of all corrective and improvement action items, as a result of non-conformities raised following the assessment.
  - Support the Regional/Country Quality Team in fulfilling the objectives of the QA function.
  - Maintains, review and updates Site Manual.
  - Manage and maintain CAPA report system includes trending and reporting
- L. QA/QC activities relating to production**
  - Review production tickets to ensure accuracy before it is released into production.
  - Review and approve redressing related work instructions before it is sent to customer for acknowledgement.
  - Perform final QA release of product to market for DHL, reject if required.



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- Support QC activities e.g. inbound QC check, in-process QC Check, final QC check, if required, including approval and rejection of products or packaging materials.
- Supervise QC staff in redressing operation.
- Review and verifies deviation reports

### *M. Support to Operations*

- Lead and provide guidance to the operations team on quality related matters.
- Attends OPS Meeting with the customer.
- Ensure RCA is performed to determine the root cause in CAPA responses
- Ensure CAPAs are responded/closed on time
- Drive and ensure monthly Management Review is conducted.
- Respond to customers quality related issues in the warehouse
- Perform other functions assigned by LSHC QA Manager and/or LSHC Director

## **5) QUALITY ASSURANCE OFFICER**

***DHL Supply Chain Singapore Pte Ltd*** (Department: **Healthcare Hub**)

7 Changi South Street 2, #01-00 Xilin Districentre Singapore 486415

*Inclusive Date: 25 Feb 2011 to 30 Apr 2013*

### Job Description/Responsibilities:

- Assists QA & Regulatory Pharmacist to oversee site quality management system and quality initiatives
- Assists QA & Regulatory Pharmacist to compile data on monthly quality/regulatory, process and quality related issues
- Assists QA & Regulatory Pharmacist to ensure that document control of quality related documents are maintained timely and effectively
- Approve or reject secondary operation materials, packaging materials and finished product
- Conduct in-process check for redressing/secondary operation, kitting operation, etc.
- Assist QA & Regulatory Pharmacist to analyze the production ticket after each production and do reconciliation, provide closure for any deviation
- Supervise QC staff in the repackaging/redressing operation
- Conduct SOP training of staff
- Assist in conducting timely checks on status of inbound pending cargo and outbound shipment on hold cargo.
- Assist QA & Regulatory Pharmacist in the management and maintenance of CAPA Report System.
- Liaise with customer on work instructions and deviation reports.
- Writes & revise SOPs and Work Instructions.
- Conduct internal audit/self-inspection

## **6) PHARMACY TECHNICIAN**

***Jurong General Hospital (Alexandra Hospital)***, Singapore

*Inclusive Date: 24 May 2010 to 24 February 2011*

### Job Description/Responsibilities:

- Assists the Pharmacists in preparing, dispensing and inventory management of pharmaceuticals and housekeeping
- Interact with patients, caregivers and other healthcare providers to achieve desired patient outcome
- Give counseling to patients and do bedside dispensing
- Help collate and monitor pharmaceutical data and reports
- Designated as Inventory & Wastage In-Charge
- Designated as in-charge of Emergency Ordering
- Do cashiering as need arises



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### **7) PHARMACY TECHNICIAN**

**Guardian Health & Beauty**, Singapore

*Inclusive Date: 23 October 2009 to 20 May 2010*

#### Job Description/Responsibilities:

- Strict adherence to Guardian Webcam Pharmacy System Standard Operation Procedures and Guardian Good Practice Guidelines
- Assist Pharmacist in dispensing medications
- Ensure accuracy of the medicine quantities packed and dispensed
- Daily back-up of files stored in hard disk
- Labels and pre-packed medicines received
- Always keep the dispensary area clean and sanitized, and medicines are arranged in proper and suitable places
- Do inventory and expiry check regularly
- Do basic counseling to customers, advises on the effective use of medicines and other health products, health issues and lifestyle choices
- Assist customers and provide information about the over-the-counter medicines and make recommendations when necessary
- Monitor and record fridge temperature.

### **8) RETAIL EXECUTIVE**

**Guardian Health & Beauty (IMM Branch)**, Singapore

*Inclusive Date: 20 April 2009 to 23 October 2009*

#### Job Description/Responsibilities:

- Assist the Store Manager in the day-to-day store operations
- AsShift-in-charge whenever the Store Manager is not around
- Do merchandising, execute promotions, planograms based on the weekly bulletin
- Do receiving of goods/stocks
- Do weekly expiry check
- Maintains cleanliness and orderliness of the shop floor
- Do cashiering as need arises
- Assist customers in their needs
- Assist Pharmacist as need arises

### **9) FOOD AND DRUG REGULATION OFFICER II**

**Department of Health (DOH) - Center for Health Development Bicol**, Provincial DOH Office, Virac, Catanduanes, Philippines

*Inclusive Date: 28 December 2000 to 10 April 2009*

*Additional Function: **Designated Officer-In-Charge Provincial DOH Representative (Team Lead)***

*Inclusive Date: 01 June 2006 to 10 April 2009*

#### Job Description/Responsibilities as Food & Drug Regulation Officer II:

- Conduct inspection of food, drug, cosmetics, medical device and household hazardous substances establishments, food manufacturers/processors to ensure compliance to DOH-BFAD regulations and ASEAN Directive
- As cGMP Auditor, performs GMP quality audits
- Prepares/writes Standard Operating Procedures (SOPs)
- Evaluate applications to ensure that all regulated products to be marketed and all regulated establishments meet the required standards, of safety, efficacy and quality
- Recommend issuance of License to Operate, Cease and Desist Order, Temporary Closure Order and Warnings to food, drug, cosmetics, medical device and household hazardous substances establishments; Food manufacturers/processors



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- Recommends Corrective and Preventive Action (CAPA) to regulated establishments
- Monitor products/ Post Market Surveillance in the markets, confiscates and condemns violative products
- Collects sample for random testing
- Conduct special investigations as directed by the BFAD and DOH-CHD Director
- Carry out investigations of complaints and handle conciliation/mediation conferences on consumer complaints
- Acts as resource person/speaker during meetings/seminars/conferences and appear in television & radio for information dissemination
- Coordinates with government agencies, Non-Government Offices (NGO), and other stakeholders for the implementation of rules and regulations of mutual concerns
- Perform other functions assigned by Regional Director.

### **Job Description/Responsibilities as OIC-Provincial DOH Representative:**

#### **A. Local Health System:**

- Coordinates/participates to Health Board Meetings
- Disseminates national policies/recommends issues for policy development
- Assists Local Government Units (LGU) on preparation of health plans.

#### **B. Public Health Reform:**

- Assists in Capability Building/Advocacy of DOH program in thrusts and priorities
- Facilitates establishment of partnership alliance with other health sectors
- Facilitates monitoring and evaluation of health program and projects.

#### **C. Hospital Reforms:**

- Facilitates extension of logistic support to hospital.

#### **D. Health Financing:**

- Advocates Health Insurance enrollment by LGUs

#### **E. Health Regulation:**

- Advocates compliance to health standards and policies
- Reviews, consolidates Technical Assistance Project and facilities provision of Technical assistance to LGUs

#### **F. Administrative:**

- Conducts monthly conferences
- Consolidates, analyzes and submits reports
- Supervises eight (8) PHT personnel (Nurses, Medical Technologist and Administrative Staff);
- Perform Performance Appraisals of personnel

### **10) HOSPITAL PHARMACIST III**

***Juan M. Alberto Memorial District Hospital***, San Andres, Catanduanes, Philippines

*Inclusive Date: 01 June 1996 to 27 December 2000*

#### **Job Description/Responsibilities:**

- Dispenses prescribed medicines for in-patients and out-patients
- Advises patients regarding the proper use of drugs, its dosage effects and other drug information
- Check patients for discharge, totals their bills for discharge and clearance
- Records dispensed medicines
- Records number of narcotic medicines issued
- Records prescriptions on the book of issues
- Prepares monthly reports on prescribed medicines
- Requisition medicines from the storekeeper
- Labels all drug container issued
- Maintains control and security of all drugs including dispensing, record keeping and secured storage of narcotics and controlled medications

### **11) RETAIL PHARMACIST**

**Catanduanes Pharmacy**, San Isidro Village, Virac, Catanduanes, Philippines

*Inclusive Date : 01 October 1995 to 31 May 1996*

#### **Job Description/Responsibilities:**



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- Prepares, dispenses and calculates the cost of prescription drugs properly prescribed by physicians and other health practitioners
- Do cashiering as need arises
- Maintains records and reports information of all prohibited and non-prohibited drugs in compliance with all state and local laws, regulations and standards regarding the handling and dispensing of drugs and medications
- Advises customers on the effective use of medicines and other health products, health issues and lifestyle choices
- Provide information about the over-the-counter drugs and make recommendations after asking series of health questions such as whether the customer is taking any other medications
- Strictly complies with the laws and regulations governing the distribution of prohibited drugs and poisonous drugs
- Always keep the Prescription section clean and sanitized and medicines are arranged in proper and suitable places
- Checks labeling requirements of stocks regularly for expiring batch, maintains proper storage condition especially for antibiotics, parenteral and vitamin preparations.

### **EDUCATION:**

- ❖ *Post Graduate Course:*  
**Master in Public Administration (MPA)**, May 2008 (6 Academic Units)  
Catanduanes State University, Virac, Catanduanes, Philippines
- ❖ *Tertiary:*  
**Bachelor of Science in Pharmacy**, 1991 – 1995  
University of Santo Tomas, Manila, Philippines

### **TRAININGS / SEMINARS / COURSES ATTENDED:**

- *Will be provided upon request*

### **REFERENCES:**

- 1. Mr. Ng Yun**  
*Pharmacist (Former Supervisor)*  
Singapore  
Mobile No.: (+65) 9152 9795
- 2. Mr. Linson Nazareno**  
*Pharmacist (Former colleague)*  
Singapore  
Mobile No.: (+65) 8222 2365

### **PERSONAL DETAILS:**

*Date of Birth:* 14 Sep 1974

*Place of Birth:* Legazpi City,  
Philippines

*Citizenship:* Filipino

*Religion:* Roman Catholic

*Civil Status:* Married

*Dependents:*

*Spouse:* Julie Ann T. Bagadiong

*Children:*

Leoj Myro T. Bagadiong

Liandel T. Bagadiong

Mari Annreia T. Bagadiong

*Address:* Blk. 335 Clementi Avenue  
2 #06-16 Singapore 120335

*Overseas Address:* San Roque, Virac,  
Catanduanes, Philippines

**Notice Period:** 1 month