CURRICULUM VITATE

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CAREER OBJECTIVE:	To associate my self with an organization that assists me to work on my proficiencies and improve my knowledge in industry. I should like to be a part of the team that works dynamically towards the growth of the organization and myself symbiotically. Looking for a challenging and responsible position to excel in the field of Pharmaceuticals with strong background in Analytical Development and Quality Control for about 14 years.
EDUCATIONAL DETAILS:	**M.Sc. (Chemistry)** from Vinayaka Missions University. **B.Sc.** Chemistry, Maths & Physics from Berhampur University, Orissa.
IT SKILLS:	• LIMS system(Caliber LIMS and star LIMS) • Nugenesis system and Track wise system • Cara Documentation and Smarteam documentation • Expertise in Windows (XP, 2010, 2013 etc Operating System and Office tools. • Ms Word, MS Excel, Power Point MS Projects. • (Word, Excel, Power Point), Data Structures.
PROFFESSIONAL PROFILE:	1 Validation of Analytical Methods like Assay by HPLC, Related Substances by HPLC, Residual solvents by GC, Genotoxic impurities by GC-MS and Cleaning methods by UV-Visible Spectrophotometer according to the Regulatory requirements. 2 Analytical method development by GC for in process, Intermediate, Raw material, Recovery solvents and drug substances. 3 Analytical method development by GC-MS for Genotoxic impurities in drug substances. 4 Analytical method Validation by GC-MS and LC-MS for Genotoxic impurities in drug substances. 5 Unknown peaks identification, impurities standard qualification and structural elucidation by by GC-MS. 6 Preparation and reviewing of Analytical Method Validation protocols and Reports as per the Regulatory requirements. 7 Evaluation and trouble shooting of problems aroused during the Method Validations in the Plant QC. 8 Reviewing and Participation in Analytical Method Transfers from ARD to the Plant QC. 9 Reviewing periodic Calibration / Performance check of Analytical Instruments. 10 Responsible and planning executed for Stability studies for Drug Substances as per the ICH/USFDA/EDQM Guidelines. 11 Preparation and review of SOPs, Validation Protocols, reports and implementation of Quality Systems as per the cGMP and GLP requirements. 12 Co-coordinating during customer audits and regulatory audits.

	13 Reviewing and planning execution of finished product, raw materials and intermediates. 14 SOP and Guidelines skilled training given to all ADL concerns as per matrix. 15 Preparation and review of Method of analysis, SOP and SOP for ADL and AVL department. 16 Periodical review of Audit trail data in LCMS and GC-MS software. 17 Reviewing, planning and execution of Mass analysis data in day to day activities.

ACHIEVEMENTS:	• Successfully completed, **Technology transfer, method validation** of various API molecules. • Customer audit faces DRL, SUNPHARAMA, GSK, TORRENT, etc. • Faced MHRA audit in March 2007 • Faced USFDA audit in May 2006. • Faced Glaxo Smithkline customer audit in Feb 2008 • Faced **WHO GENEEVA** in year 2008 as a Sr.Officer for Three products at a time without any non compliance. • Faced KFDA audit in Feb-2011. • Supporting to resolve regulatory queries.
STRATEGIC PLANNING:	• Steering operations with a view to achieve organizational objectives and ensure profitability. • Formulating and materializing monthly / annual goals, short / long term budgets and developing Operational plans for the achievement of these goals. • Devising and implementing policies & procedures to enable smooth functioning of operations. • Good knowledge in analytical method validations by using HPLC, GC and other techniques for API's.
Technical Competencies	Operation and Calibration of the following Instruments : o HPLC – Agilent (1200 Series with Ezchrome &Chemstation), Waters (Alliance with 2487 and 2996(PDA) Detector with Empower software, Shimadzu Class VP with LC-Solution software. o GC – Agilent 6890, 7890 and 7890B with Empower software o GC – Perkin Elmer with Empower and Total chrome software o GCMS- Agilent GCMS 7000,7010,7010B and 5975Cwith MSDCHEM and Mass hunter software. LC-MS- Triple quadrupole (TQ)-Make-Water with Masslynx software. o FT IR – Thermo Nicolet, Shimadzu o UV-VIS Spectrophotometer – Shimadzu o KF & Autotitrator – Metrohm. o Qualification of all Analytical Instruments & equipments. o Documentation and review of IQ,OQ and PQ related documents.

	o Raising and Closing of Change Controls and Deviations and handling of Out-Of-Specifications and OOT. o Experienced in implementation and auditing of Quality Systems as per the requirements of regulatory agencies. o Effective Time Management for completion of projects by planning with excellent Team co-ordination and support. • Excellent strengths in Team management, oral, written and communication skills. • Facing so many regulatory audits like WHO Geneva, CGMP, and MHRA & USFDA. • Controlling of manpower management in Analytical development lab. • Coordinating with QA and RA department for DMF filling in various products. • Find the Literature and technical data support for analytical method development.
COMMERCIAL FUNCTIONS:	• Assessing annual requirement of material in line with core organizational objectives. • Interfacing with key influencers among corporate for assessing requirements and delivering need based product solutions. • Reducing solvent and reagent conservations. • Indention and following up the indent material with purchase department.
QUALITY:	• Interacting with the CRD people for daily planning and routine analysis support. • Interacting with the CRD people for freeze the specification finalize the STP. • Maintaining GLP Systems and process Control Records. • Assuring conformance of the product to safety norms and specifications. • Ensuring maintenance of the proper Working Conditions, Housekeeping, Discipline and Safety at workstations.
PEOPLE MANAGEMENT:	• Coordinating with All relevant departments like: ware house, production, CRD, ARD, RA, QC,QA, HR and Safety in conducting negotiations with towards resolving issues / grievances, signing agreements and maintaining amicable employee relations. • Overseeing the performance of subordinates, motivating them to improve their contribution levels. • Organizing Skill Development Training along with the HOD-HR and Training with QA Depts.

WORK EXPERIENCE:	• Presently working in Mylan Laboratories, LTD as a Manager in Analytical development Laboratory (ADL) department since Sept-2013. • Worked in TEVA API INDIA LTD, GAJRAULA as an Officer-III in Analytical Technology Services (ATS) department from May-2012 to Aug-2013. • Worked in SURYA PHARMACEUTICALS LTD. BANUR as a Research Scientist in Analytical Research & Development (ARD) department from March-2010 to May-2012. • Worked in DHARUHERA PHARMACEUTICALS LTD as an Executive in Analytical Research & Development (ARD) department from Aug-2009 to March-2010. • Worked in MACLEOD'S PHARMACEUTICALS LTD as a Sr.officer in Analytical Method Validation (AMV) department from Sept-2007 to July-2009. • Worked as a Chemist in AUROBINDO PHARMA LTD, UNIT-XI, from March-2006 to Sept-2007.
TOTAL EXEPERIENCE:	• 14 years

INTERESTS:	• Playing and watching Cricket. • Exchanging Knowledge with friends. • Reading Magazines & Books, Enjoying Music & Traveling, Eagerness to know more & more.
STRENGTHS:	• Hard working nature. • Optimistic and Self-Confidence. • Team facilitator.
PERSONAL DETAILS:	Name : Arata Trana Mahapatra. Father's Name : Gopal Chandra Mahapatra. Native place : Odisha Date of Birth : Ist June, 1980 Sex : Male Marital Status : Married Languages Known : English, Hindi, Oriya and Telugu
Present CTC	13.4 LACS
Notice Period	3.0 Months
DECLARATION:	I hereby declared that above given information is true to best of my knowledge and I Shall be grateful to you if you consider my candidature. I can assure you that I will do my best and dedicate my Service to your discretion.

Place: Hyderabad, Telangana.

Date: 14.09. 2019 **(Arata Trana Mahapatra)**