# CURRICULUM VITAE

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| **Full Name** | **First** | **Middle** | **Last** |
| Kinjalkumar | Chaturbhai | Patel |
| **Address** | Plot No-1469/1, Sector – 5 B, Gandhinagar - 382006 | | |
| **Birth Date** | 23-February-1990 | | |
| **Sex** | Male | | |
| **Hobby** | Reading, Listening Music | | |
| **Nationality** | Indian | | |
| **Telephone Number** | +91-8735983508 | | |
| **Email id** | [Patelkccology@gmail.com](mailto:Patelkccology@gmail.com) | | |

**Education/Academic Qualifications:**

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| **Degree** | **Year** | **Institution, Country** | **Result** |
| M.Pharm | 2013 | Shree S.K. Patel College of Pharmaceutical Education and Research, Ganpat University, Kherva, Mehsana,  Gujarat, India | 8.34 CGPA |
| B.Pharm | 2011 | Kalol Institute of Pharmacy, Kalol, Gandhinagar, Gujarat, India | 72.25% |
| H.S.C | 2007 | Shree Swaminarayan Higher Secondary School, Gandhinagar, Gujarat, India | 86.40% |
| S.S.C | 2005 | Shree Swaminarayan High Secondary School, Gandhinagar, Gujarat, India | 85.14% |

**Employment Experience:**

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| **Start and End Dates** | **Title** | **Institution or Company, State/Province/Country along with brief description of Job Responsibilities** |
| 15-Nov-2018 – 30-Jan-2021 | Senior Research Fellow - Clinical Research - Pharmacovigilance | Emcure Pharmaceuticals Limited  Uvarsad Square, Sarkhej Gandhinagar Highway, Adalaj, Gandhinagar – 382421  **Job Description:**   * Mentor of Triaging activity * Give training to employee to do Literature activity, regulatory cases triaging. * Process ICSRs when needed and submit them to different regulatory authority when required. * Aggregate Reporting: * Preparation and reporting of different aggregate reports (eg-PADERs, PBRERs, PSURs Addendum to clinical overview summary report) and Risk Management plan as per company controlled documents (e.g. SOPs, WIs, Guide) and guidelines. * Responsible for management and timely submission of individual case reports as well as aggregate report in order to achieve 100% regulatory compliance. * Signal Detection: * Performing signal detection and management activities for company products as well as contribute in performing benefit risk analysis of the product when needed. * Contributing to the on-going enhancement of Pharmacovigilance processes and preparing standard operating procedures related to pharmacovigilance, as needed. * Providing support as needed for regulatory authority inspections and conduct training to new employees. |
| 06 Feb 2017 to 06  Nov 2018 | Senior Research Associate – Clinical Research - Pharmacovigilance | **Lambda Therapeutic Research Ltd, Ahmedabad**  Job responsibilities- **Literature review process**. **Case processing** and reporting the ICSRs to the various regulatory authorities within time frame and entering or updating the drug information on **xEVMPD** for various marketing authorization holders.  Review of the Regulatory Response(USFDA, EU, Canada and WHO)  Preparation of SOP’s, Forms, Templates and Checklists.  Give training to other person for Literature review and Submission. |

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| 01 Jun 2015 to 30  Jul 2016 | Medical Record Reviewer – Coding Department | **Advantmed Solutions, Ahmedabad**  Job responsibilities- working on patients all details like its past history, current diagnosis, surgical procedures and medication and do coding for all process as per ICD 10 guideline and provide coding detail to Insurance  company for their claiming procedure |
| 07 Nov 2011 to 30  Apr 2015 | Jr. Pharmacist | Shree Jay Ambe Medical Stores  Job responsibilities- Dispensing of medicines |

**Current Job:** From 03-Feb-2021 to till date

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| **Present position** | Senior Executive |
| **Department** | Pharmacovigilance |
| **Office Address** | Lambda Therapeutic Research Ltd, Ahmedabad |

# Current Job Responsibilities:

# Aggregate Reporting:

# Preparation and reporting of different aggregate reports (eg-PADERs, PBRERs, PSURs Addendum to clinical overview summary report) and Risk Management plan as per company-controlled documents (e.g. SOPs, WIs, Guide) and guidelines.

# Responsible for management and timely submission of individual case reports as well as aggregate report to achieve 100% regulatory compliance.

# Providing support as needed for regulatory authority inspections and conduct training to new employees.

# Trainings/Certifications:

* + Successfully completed B.Pharm Industrial training at Wellable Pharmaceuticals, Mehsana of 3 months.
  + ICMR sponsored two days national seminar on ‘Novel Paradigms In Reverising Diabetes And Diabetic Complications’ at Shree S.K.Patel college of Pharmaceutical education & Research, Kherva, Ganpat University.
  + DST sponsored three days national seminar on ‘Training and Hands on Molecular Biotechnology Tools and Techniques’ at Shree S.K.Patel college of Pharmaceutical education & Research, Kherva, Ganpat University.
  + 3rd National Seminar on “Quality Assurance Issues in Drug Discovery & Formulation Development” held at Babaria Institute of Pharmacy, Varnama, Vadodara
  + 29th Annual Conference of Indian Pharmacological Society – Gujarat Chapter 2013 held at Ahmedabad.

# Dissertation:

M. Pharm

* “The role of HDAC inhibitors in Diabetes and associated cardiovascular complication in Rats” (Research)
* “The Glamour and Gloom of HDAC” (Review)

# Other Details:

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| **Membership of Scientific Societies** | No Yes  If yes, specify or attach document  Member of Gujarat state pharmacy council with registered pharmacist number G45559 |
| **Publications** | No  Yes If yes, specify or attach document |

**Declaration:**

I, hereby declare that all the entries are correct to best of my knowledge. If chance is given to me I assure you that, I will prove myself as an asset to your organization.