**CURRICULUM VITAE**

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| **1. Name**: Van Thi My Hue |
| **2. Date of birth**: 13/9/1975 **3. Gender**: F |
| **4. Academic title:** PhD |
| **5. Administrative position:** Lecturer |
| **6. Address**: 25 Han thuyen, Pham Dinh Ho - Hai Ba Trung - Hanoi |
| **7. Telephone**: 84-1234266388 **E-mail**: vanmyhue2@yahoo.com |
| **8. Affiliation:**   **Institute** : Organic Chemistry Department, Hanoi University of Pharmacy, Vietnam **Address** : 13-15 Le Thanh Tong street –Hoan Kiem district- Hanoi- Vietnam |
| 9. Qualification |
| Academic Degree | Institutions | Major | Years |
| Degree of Pharmacy | Hanoi University of Pharmacy, Vietnam | Pharmacist | 1997 |
| M. S | Hanoi University of Pharmacy, Vietnam | Pharmacist | 2000 |
| Ph. D. | Chonnam National University, Korea | Pharmaceutical Chemistry | 2008 |
| 10. Professional Experiences |
| *Year* | *Position* | ***Affiliation*** | ***Address*** |
| 2001-2004 | Lecturer | Organic Chemistry Department, Hanoi University of Pharmacy | 13-15 Le Thanh Tong –Hoan Kiem - Hanoi- Vietnam |
| 2004-2008 | Ph.D course | Chonnam National University, Korea | 300 Yongbong-dong, Buk-gu, Gwangju Korea |
| 2009-2010 | Lecturer | Organic Chemistry Department, Hanoi University of Pharmacy | 13-15 Le Thanh Tong –Hoan Kiem- Hanoi- Vietnam |
| 2010-now | Deputy Head of Organic Chemistry Department, Lecturer | Organic Chemistry Department, Hanoi University of Pharmacy | 13-15 Le Thanh Tong –Hoan Kiem- Hanoi- Vietnam |
| 11. Teaching Experiences |
| Year | Position | Class |
| 2001-2004, 2009-now | Lecturer | Organic Chemistry Department, Hanoi University of Pharmacy |
| June, 2013 ~ Sep, 2013 | Lecturer  | Master class in Faculty of Pharmacy, Mahidol University, Thailand |
| 12. Language skills/ Proficiency• Fluent in English (dissertation of PhD courses written in English) |
| **13. List of International Publications**1/**.** SAR Based Design of Nicotinamides as a Novel Class of Androgen Receptor Antagonists for Prostate Cancer. ***Journal of Medicinal Chemistry***- Brief article, **2013**. 2/. Novel hydroxamic acids having histondeacetylase inhibiting activity and anticancer composition comprising the same as an active ingredient, ***Korean Patent No. 10-2012-0066831.*** 3/.Benzothiazole-containing hydroxamic acids as histone deacetylase inhibitors and antitumor agents. ***Bioorganic and Medicinal Chemistry Letters***. **2011,**21(24), 7509-7512. 4/. Novel Hydroxamic Acids Having Histone Deacetylase Inhibiting Activity and Pharmaceutical Composition for Treating Cancer Comprising the Same As Active Ingredient. ***Korean Patent No. 10-2011-0050864*** 5/**.** Synthesis of benzo[3,4]azepino[1,2-*b*]isoquinolin-9-ones from 3-arylisoquinolines via ring closing metathesis and evaluation of topoisomerase I inhibitory activity, cytotoxicity and docking study, ***Bioorg. Med. Chem.,*** (**2011**), 19 (18), 5311-5320. 6/. Application of Ring-Closing Metathesis for the synthesis of Benzo[3,4]azepino[1,2-b]isoquinolin-9-ones, ***Chem. Pharm. Bull.***, (**2011**), 59 (9), 1169-1173. 7/. Design, synthesis of 4-amino-2-phenylquinazolines as novel topoisomerase I inhibitors with molecular modeling, ***Bioorg. Med. Chem.,*** (**2011**), 19, 4399-4404. 8/. Design, synthesis and docking study of 5-amino substituted indeno[1,2-c]isoquinolines as novel topoisomerase I inhibitors, ***Bioorg. Med. Chem.,*** (**2011**), 19 (6), 1924-1929. 9/**.** Development of 3-aryl-1-isoquinolinamines as potent antitumor agents based on CoMFA, ***Eur. J. Med. Chem***. (**2010**), 45 (11), 5493-5497. 10/**.**  Synthesis, in vitro and in vivo evaluation of 3-arylisoquinolinamines as potent antitumor agents, ***Bioorg. Med. Chem. Lett.***, (**2010**), 20 (17), 5277-5281. 11/. Total synthesis of 8-oxypseudopalmatine and 8-oxypseudoberberine via Ring Closing Metathesis, ***Tetrahedron,*** (**2009**), 65, 10142-10148.  12/. Molecular design, synthesis and docking study of benz[b]oxepines and 12-oxobenzo[c]phenanthridinones as topoisomerase I inhibitors, ***Bioorg. Med. Chem. Lett.,*** (**2009**), 19 (9), 2444-2447. 13/**.** Structural modification of 3-arylisoquinolines to isoindolo[2,1-b]isoquinolinones for the development of novel topoisomerase I inhibitors with molecular docking study, ***Bioorg. Med. Chem. Lett.,*** (**2009**), 19 (9), 2551-2554.14/. Application of Coupling Reaction between Lithiated Toluamide and Benzonitrile for the Synthesis of Phenolic Benzo[c]phenanthridine Alkaloid, Oxyterihanine, ***Arch. Pharm. Res***.,(**2008**), 31 (1), 6-9. 15/. Convenient synthesis of indeno[1,2-c]isoquinolines as constrained forms of 3-arylisoquinolines and docking study of a topoisomerase 1 inhibitor into DNA-topoisomerase 1 Complex, ***Bioorg. & Med. Chem. Lett.,*** (**2007**), 17, 5763-5767. 16/**.** Design, Docking and Synthesis of Novel Indeno[1,2-c]isoquinolines for the Development of Antitumor Agents as Topoisomerase 1 Inhibitors, ***Bioorg. & Med. Chem. Lett.,*** (**2007**), 17 (13), 3531-3534. |