# **CURRICULUM VITAE**

### Ponnappa.K.C

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## **EDUCATION:**

1999-2005. **Ph.D. Biochemistry.** Department of studies in Biochemistry, University of Mysore, India. (Thesis will be submitted in May 2006)

<u>Doctoral Thesis</u>: "Comparative characterization of the representative acidic and basic PLA<sub>2</sub> from *Naja naja naja* venom."

1993-1995. **M.Sc. Biochemistry**, Department of studies in Biochemistry, University of Mysore, India.

1990-1993. **B.Sc. Biochemistry,** Botany and Microbiology, University of Mysore, India.

### **RESEARCH EXPERIENCE:**

- 1. 1999-Till date: Working as a Resarch Fellow in the Department of Studies in Biochemistry, University of Mysore, India, for the Ph.D, in Biochemistry with the title "Comparative characterization of the representative acidic and basic PLA<sub>2</sub> from Naja naja naja venom". Work involved isolation and purification of PLA<sub>2</sub>s from Indian cobra venom. PLA<sub>2</sub>s from snake venom show high degree and homology and hence separation of individual PLA<sub>2</sub> poses a difficult task. This work has thus, made me confident in designing and carrying out experiments, enabled me to develop my skills in isolation procedures and handling animals. This has given me opportunity to work in collaboration in different laboratories. The details of study carried out,
- **Purification and characterization** of snake venom enzymes (such as PLA<sub>2</sub>s, Three finger toxins and PLA<sub>2</sub> inhibitor.) by using various purification techniques like Column chromatography (gel filtration as well as ion exchange chromatography) and RP-HPLC

- N terminal sequencing of purified proteins; Acidic PLA<sub>2</sub> and cyto/cardiotoxin
- Interaction studies with inhibitors (Aristalochic acid and endogenous PLA<sub>2</sub> inhibitor NN Ic from Indian cobra venom.) and studying the nature of inhibition by employing both biochemical as well as biophysical parameters such asIC<sub>50</sub>, enzyme inhibition constant k<sub>I</sub>, and interaction monitoring by following fluorescence quenching
- **Protein modification studies** were carried out to determine the active site residues by modifying histidine residue using pBPB (bromophenacylbromide) in PLA<sub>2</sub>s. In modifying the tryptophane residue in three finger toxin to study the role of it in lethality.
- **Pharmacological activities** like PLA<sub>2</sub> activity, neurotoxicity, myotoxicity, cytotoxicity, edema inducing activity, proteolytic activity, hemorrhagic activity
- Platelet aggregation activity inhibition was tested for acidic D IV PLA<sub>2</sub> and aggregate Ag III PLA<sub>2</sub>. Effect of aspirin on platelet aggregation
- **Toxicity assays** including, LD<sub>50</sub>, isolation of sciatic nerve gastronemius muscle from frogs were carried out to determine the mode of toxin interaction.
- Raising and purifying antibody from chick. The effect of IgY prepared against the venom and
  purified toxin on the pharmacological properties and cross reactivity and neutralization of venom
  toxicity, efficacy of IgY.
- **Animal experiments**: Experience in handling mice, rats, frogs and rabbits. Determination of LD<sub>50</sub>, myotoxicity, neurotoxicity and hemorrhagic activity using mice. Mouse paw model for edema inducing activity.

# **MAJOR FINDINGS OF MY RESEARCH**:

- Isolation and characterization of unique toxic acidic PLA<sub>2</sub> enzyme from cobra (*Naja naja naja*) venom, its effect on platelet aggregation activity, its interaction with inhibitors and N terminal sequence.
- Isolation of hetero trimeric NNXI (PLA<sub>2</sub>) a post synaptic complex a first report .Its contribution to the venom toxicity.
- First report of unusual behavior of PLA<sub>2</sub>s upon separation of basic PLA<sub>2</sub>s from the peptide pool. Report of a artificially formed PLA<sub>2</sub> complex devoid of pharmacological activities
- Isolation and characterization of three finger toxins from *Naja naja naja* venom .Interaction of cardiotoxin with different cell line (EAT and leukocytes), chemical modification leading to loss of lethality. First report of non interaction of cardiotoxin with leukocytes indicating the presence of recognition site for cardiotoxin action. Also N terminal sequence done to show its uniqueness.
- Raising and purifying antibody from chick. The effect of IgY prepared against the venom and purified toxin on the pharmacological properties and cross reactivity and neutralization of venom toxicity, efficacy of IgY for possible therapeutic use.
- 2. Project work carried out during Masters Degree (Biochemistry) Course (1993-1995) entitled "Partial purification of a PLA<sub>2</sub> from Vipera russelii venom". This work first exposed me to the world of proteins. I had partially purified a PLA<sub>2</sub> from the venom of Vipera russelii, by chromatographic techniques such as gel filtration and ion exchange chromatography. Followed by PAGE show its partial purification from the venom.

## **TEACHING EXPERIENCE AND OTHER ASSIGNMENTS:**

- Teaching assignment carried out (from 1999 to date) along with research work for postgraduate and graduate students in both chemistry and biochemistry. I have taught bioorganic chemistry, Biochemical Techniques, Nutrition, Protein Biochemistry, Enzymology, Metabolism, Cell biology and Molecular biology.
- 2. Experienced in guiding more than 20, M.Sc., students for their project work during my research period in different topics of biochemistry.

# **PUBLICATIONS:**

- C.T.Sadashiva, J.N.Narendra Sharat Chandra, K.C.Ponnappa T.Veerabasappa Gowda and K.S.Rangappa Synthesis and efficacy of 1-[bis(4-fluorophenyl)-methyl] piperazine derivatives for acetylcholine esterase inhibition, as a stimulant of central cholinergic neurotransmission in Alzheimer's disease. (Accepted for publication in Bioorganic and Medicinal Chemistry Letters on 27/04/06)
- **Ponnappa.K.**C, Krishnakantha.T.P, Kini.R.M and Veerabasappa Gowda A novel toxic acidic PLA<sub>2</sub> from the venom of *Naja naja naja: isolation*, purification and Charecterization. (Communicated to Biochimie)
- Basavarajappa B.S; Ponnappa K.C; Veerabasappa gowda.T Isolation and characterization of a
  naturally occurring novel α toxin composed of PLA<sub>2</sub> isomers from Indian cobra Naja naja
  venom. (Communicated to Toxicon)
- **K.C.Ponnappa** and T.Veerabasappa Gowda The contribution of peptides and low molecular weight components in the venom towards PLA<sub>2</sub> stability and lethality of south Indian cobra (*Naja naja naja*) venom. (Manuscript under preparation)

### **BOOK CHAPTERS:**

• "Indian Cobra *Naja naja naja* venom:Composition enzymes and toxins". **K.C.Ponnappa** and T.Veerabasappa Gowda -a full length chapter for the book "**The great mystic cobra**" Edited by Dr N.S.Leela, Indian Institute of Science (to be published shortly)

## **TECHNICAL EXPERTISE:**

**Protein purification** : Gel permeation, Ion exchange and affinity

Chromatography, TLC, HPLC and RP- HPLC.

**Protein analysis** : Acrylamide gel electrophoresis, Western blotting.

Spectroscopy and Spectrofluorophotometry.

**Radio labeling** : Can handle radioactive material and label *E.coli* and other

cells with radioactive chemicals.

**Immunological techniques**: Immunization and development of antibodies in Rabbit,

development of Hen's egg yolk IgY antibodies. ELISA and

Immunodiffusion.

**Microbiological techniques**: Preparation of culture media (liquid and agar plates),

Isolation of pure culture, Identification of bacteria by staining

methods & sensitivity analysis of biological samples.

**Animal experiments**: Experience in handling mice, rats and rabbits.

Determination of LD<sub>50</sub>, myotoxicity, & hemorrhagic

activity using mice. Mouse paw model for edema inducing

activity. Antibody (polyclonal) production using chick and

Rabbit as animal model.

**Enzyme kinetics**: PLA<sub>2</sub> assay, Protease assay, Fibrin (ogen) olytic assay,

Hyaluronidase assay, Gelatinolytic assay (Zymogram),

Acetylcholine esteraease assay, Procoagulant and anticoagulant assays, their kinetic & interaction studies

with activators and inhibitors.

**Modification studies of Enzyme** 

**Catalytic sites** 

: Chemical modification of amino acid residues of histidine

and tryptophan

Other qualifications : Good computing experience in Word, Excel, Power point

and Image processing. Sequence Homology by BLAST

search

## **HONORS AND AWARDS:**

1. **Recipient of Senior Research fellowships** from the University Grants Commission, Government of INDIA.

- 2. **Recipient of Junior Research fellowships** from the University Grants Commission, Government of INDIA
- 3. **Eligible for lectureship** by passing State Level Entrance Test conducted by Government of Karnataka, INDIA

#### Participation in scientific meetings and symposiums:

- 1) National symposium on Proteins in Biochemistry: Structure, Function, Expression and specificity of action organized by Biochemistry Alumni Association, University of Mysore, India. November 15 –16 (1999).
- 2) National Symposium on "Control of Animal experiments through Alternative approaches" Organized by CPCSEA, Ministry of Social Justice and Empowerment .Government of India. Defence Food Research Laboratory, Mysore, India. March 25 (2000).
- 3) "Contribution of PLA<sub>2</sub> towards the lethal toxicity of Indian cobra venom" Ponnappa.K.C and Veerabasappa Gowda presented at 3rd National Symposium on Venoms and Toxins, organized by Biochemistry Alumni Association, University of Mysore, Mysore, India. December 23-24, 2003.

#### MEMBER TO SCIENTIFIC BODIES:

- ♦ Society of Biological Chemists, India.
- ♦ Society of Applied Botany and Biotechnology, India.
- ♦ Association of Carbohydrate Chemist and Technologist, India.

# **PERSONAL**:

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**Date of Expiry** : 24/02/2008

Place of Issue : Bangalore, India.

# **REFERENCES:**

## 1. Prof.T.Veerabasappa Gowda. (Ph.D supervisor)

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