# **CURRICULUM VITAE**

## PERSONAL DATA:

Name:

Yanfen Hu

Date of Birth:

December 13, 1963

Place of Birth:

Shanghai, China

Citizenship:

United States of America

Marital Status:

Married, two children

Social Security Number: 113-70-6225

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# **EDUCATION AND POST-GRADUATE TRAINING:**

1981-1985	Fudan University, China	B.S. in Genetics
1987-1992	University of California, Berkeley	Ph.D. in Molecular Biology
1992-1994	Genentech, Inc., California	Postdoc. in Apoptosis
1994-1996	Cold Spring Harbor Laboratory	Postdoc. in Signal Transduction

## ACADEMIC APPOINTMENTS:

Research Associate 1996-2002

Department of Biochemistry and Molecular Genetics

University of Virginia

Assistant Professor of Research 2002-2005

Department of Biochemistry and Molecular Genetics

University of Virginia

Associate Professor of Research 2005-present

Department of Biochemistry and Molecular Genetics

University of Virginia

#### HONORS AND AWARDS:

1986-1987 China-US Biochemistry Examination and Application (CUSBEA) Scholarship;

Total score ranked the third place and Molecular Biology subject ranked the first

place nationwide

1994-1995 Robertson Fellowship, Cold Spring Harbor Laboratory

## PROFESSIONAL AFFILIATIONS:

American Association for Advancement of Science (AAAS) American Association of Cancer Research (AACR) The Endocrine Society

#### AREAS OF RESEARCH INTEREST:

Breast cancer development and prevention. Gene expression and regulation.

#### **RESEARCH FUNDING:**

### Current grant support:

5/03-4/06 Department of Defense (DAMD17-03-1-0398, P.I.: Yanfen Hu)

Cooperation between BRCA1 and JunB in Gene Regulation and Tumor Suppression

Direct cost per year: \$100,000

## Pending research grant application:

12/05-11/10 NIH (RO1 CA118578, P.I. Yanfen Hu)

The Mechanism of BRCA1 in Tumor Suppression

Direct cost per year requested: \$200,000

15.6% after first submission as a New Investigator

1/06-12/08 Department of Defense (BC050500, P.I. Yanfen Hu)

The Function of Putative Tumor Suppressors BRCA2 and BARD1 in Aromatase

Expression and Estrogen Biosynthesis.

Direct cost per year requested: \$100,000

### **PUBLICATIONS:**

- 1. Orias, E., Larson, D., **Hu, Y-F.**, Yu, G., Karttumen, J., Lovlie, A., Haller, B., and Blackburn, E.H. (1988) Replacement of the macronuclear ribosomal RNA genes of a mutant Tetrahymena using electroporation. *Gene* 70: 295-301
- 2. **Hu, Y-F.**, Luscher, B., Admon, A., Mermod, N., and Tjian, R. (1990) Transcription factor AP-4 contains multiple dimerization domains that regulate dimer specificity. *Genes & Dev.* 4: 1741-1752

- 3. Tartaglia, L.A., Rothe, M., Hu, Y-F., and Goeddel, D.V. (1993) Tumor Necrosis factor's cytotoxic activity is signaled by the p55 TNF receptor. *Cell* 73:213
- 4. **Hu, Y-F.**, Hao, Z., and Li, R. (1999) Chromatin remodeling and activation of DNA replication by the BRCT-domain of BRCA1. *Genes & Dev.* 13: 637-642
- 5. Miyake, T., **Hu, Y-F**., Yu, D.S., and Li, R. (2000) A functional comparison of BRCA1 C-terminal domains in transcription activation and chromatin remodeling. *J. Biol. Chem.* 275:40169-40173
- Hu, Y-F., Miyake, T., Ye, Q., and Li, R. (2000) Characterization of a novel trans-activation domain of BRCA1 that functions in concert with the BRCA1 C-terminal (BRCT) domain. J. Biol. Chem. 275:40910-40915
- 7. Fourel, G., Boscheron., C., Revardel, E., Lebrun, E., **Hu, Y-F**., Carmine, K., Muller, K., Li, R., Mermod, N., and Gilson, E. (2001) An activation-independent role of transcription factors in insulator function. *EMBO reports* 2:124-132
- 8. Ye, Q., **Hu**, Y-F., Belmont, A., and Li, R. (2001) BRCA1-mediated high-order chromatin unfolding and its deregulation by cancer predisposing mutations. *J. Cell Biol.* 155:911-921
- 9. **Hu, Y-F.**, and Li, R. (2002) JunB potentiates function of BRCA1 activation domain 1 (AD1) through a coiled-coil-mediated interaction. *Genes & Dev.* 16:1509-1517
- 10. Wu, Y., Ghosh, S., Nishi, Y., Yanase, T., Nawata, H., and **Hu, Y-F**. (2005) The Orphan Nuclear Receptors NURR1 and NGFI-B Modulates Aromatase Gene Expression in Ovarian Granulosa Cells: a Possible Mechanism for Repression of Aromatase Expression upon Luteinizing Hormone Surge. *Endocrinology* 146:237-246
- 11. Ghosh, S., Wu, Y., Li, R., and **Hu, Y-F.** (2005) Jun proteins modulate the ovary-specific promoter of aromatase gene in ovarian granulosa cells via a cAMP-responsive element. *Oncogene* 24:2236-2246
- 12. **Hu**, Y-F.\*, Ghosh, S., Amleh, A., Yue, W., Lu, Y-Z., Katz, A., and Li, R.\* (2005) Modulation of Aromatase Expression by BRCA1: A Possible Link to Tissue-Specific Tumor Suppression. *Oncogene* in press (\*co-corresponding authors)

### INVITED MEETING PLATFORM PRESENTATION:

International Conference on Aromatase (Edinburgh, 2004) DOD Era of Hope Breast Cancer Research Conference (2005)