

## CV



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#### Education

1988-1992 B. A., Microbiology, Seoul National University, Korea  
1992-1994 M. S., Microbiology, Seoul National University, Korea  
1994-1998 Ph.D., Biochemistry, Oxford University, UK

#### Major Activities

1999-2001 Postdoctoral Fellow, Howard Hughes Medical Institute, University of Pennsylvania, USA  
2001-2004 Research Assistant Professor, Seoul National University, Korea  
2004-2008 Assistant Professor, Seoul National University, Korea  
2008-2013 Associate Professor, Seoul National University, Korea  
2010-present Editorial Board, *Cell*  
2010-present SNU Distinguished Fellow, Seoul National University, Korea  
2011-present Editorial Board, *EMBO Journal*  
2012-present Director, Center for RNA Research, Institute for Basic Science, Korea  
2012-present Editorial Board, *Genes & Development*  
2012-present Foreign Associate Member, EMBO  
2013-present Professor, Seoul National University, Korea

#### Honors and Awards

2007 Thomson Scientific Citation Award  
2007 Young Scientist Award, Korea  
2007 Woman Scientist of the Year, Korea  
2008 L'Oreal-UNESCO Women in Science Award,  
2009 Ho-Am Prize in medicine, Korea  
2010 Amore Pacific the Grand Prize, Korea  
2010 National Honor Scientist, Korea  
2013 The Korea S&T Award, Korea

#### Research Interests

RNA-mediated gene regulation, microRNA biogenesis and function, Regulatory non-coding RNA, RNA-protein network, Stem cells and fate decision

## **Publications (Most representative 10 items)**

1. Lee Y., Ahn C., Han J., Choi H., Kim J., Yim J., Lee J., Provost P., Rådmark O., Kim S., and Kim V.N. 2003. The Nuclear RNase III Drosha Initiates MicroRNA Processing. *Nature* 425:415-419.
2. Han J., Lee Y., Yeom K.H., Nam J., Lee K., Sohn S., Cho Y., Zhang B., and Kim V.N. 2006. Molecular Basis for Primary MicroRNA Processing by the Drosha-DGCR8 complex. *Cell* 125:887-901.
3. Heo I., Joo C., Cho J., Ha M., Han J. and Kim V.N. 2008. Lin28 mediates the terminal uridylation of let-7 precursor microRNA. *Molecular Cell* 32:276-284.
4. Han J., J. S. Pederson, Kwon S. C., C. D. Belair, Kim Y. K., Yeom K. H., Yang W. Y., D. Haussler, R. Blelloch, and Kim V.N. 2009. Posttranscriptional crossregulation between Drosha and DGCR8. *Cell* 136:75-84.
5. Heo I., Joo C., Kim Y.K., Ha M., Yoon M.J., Cho J., Yeom K.H., Han J. and Kim V.N. 2009. TUT4 in Concert with Lin28 Suppresses MicroRNA Biogenesis through Pre-MicroRNA Uridylation. *Cell* 138:696-708.
6. Hyun S., Lee J. H., Jin H., Nam J.W., Namkoong B., Lee G., Chun J., and Kim V.N. 2009. Conserved microRNA miR-8/miR-200 and its target USH/FOG2 control growth by regulating PI3K. *Cell* 139:1096-1108.
7. Park J.E., Heo I., Y. Tian, D. K. Simanshu, Chang H., D. Jee, D. J. Patel and Kim V.N. 2011. Dicer recognizes the 5' end of RNA for efficient and accurate processing. *Nature* 475:201-205
8. Heo I., Ha M., Lim J., Yoon M., Park J.-E., Kwon S. C., Chang H., and Kim V. N. 2012. Monouridylation of pre-microRNA as a key step in the biogenesis of group II let-7 microRNAs. *Cell* 151: 521-532.
9. Cho J., Chang H., Kwon S. C., Kim B., Kim Y., Choe J., Ha M., Kim Y. K., and Kim V. N. 2012. LIN28A is a suppressor of ER-associated translation in embryonic stem cells. *Cell* 151: 765-777.
10. Kwon S. C., Yi H., Eichelbaum K., Föhr S., Fischer B., You K. T., Castello A., Krijgsveld J., Hentze M. W., and Kim V. N. 2013 The RNA-binding protein repertoire of embryonic stem cells. *Nature Structural and Molecular Biology*, in press.

# The RNA-Binding Protein Repertoire of Embryonic Stem Cells

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RNA-binding proteins (RBPs) play essential roles in RNA-mediated gene regulation, and yet the current annotation of RBP is largely limited to those with known RNA-binding domains. To systematically identify the RBPs of embryonic stem cells (ESCs), we here employ “interactome capture”, which combines UV-crosslinking of RBP to RNA in living cells, oligo(dT) capture, and mass spectrometry. From mouse ESCs, 555 proteins are defined to constitute the mESC mRNA-interactome, which includes 283 proteins not previously annotated as RBPs. Interestingly, 68 novel RBP candidates are highly expressed in ESCs compared to differentiated cells, implicating their roles in stem cell physiology. Among them, two well-known E3 ubiquitin ligases, Trim25 (or Efp) and Trim71 (or Lin41), are validated as RBPs, revealing a potential link between RNA biology and protein modification pathways. Our study confirms and expands the atlas of RBPs, providing a valuable resource for the study of RNA-RBP networks in stem cells.

**Keywords** : microRNA, non-coding RNA, RNA interference, RNA-binding protein