



• **Name:** Hyun-Young Kim

• **Current Position:** Clinical Assistant Professor
Department of Laboratory and Genetics, Samsung Medical Center,
Sungkyunkwan University School of Medicine, Seoul, Korea

• **Country:** Korea

• **Educational Background:**

2013 – 2015 Master of Medicine,
Sungkyunkwan University School of Medicine, Seoul, Korea

2002 – 2008 Bachelor of Medicine,
Choongnam National University School of Medicine, Daejeon, Korea

• **Professional Experience:**

Jun 2019 – present Clinical Assistant Professor,
Department of Laboratory Medicine and Genetics,
Samsung Medical Center, Seoul, Korea

Mar 2017 – May 2019 Clinical Assistant Professor,
Department of Laboratory Medicine,
Gyeongsang National University Hospital, Jinju, Korea

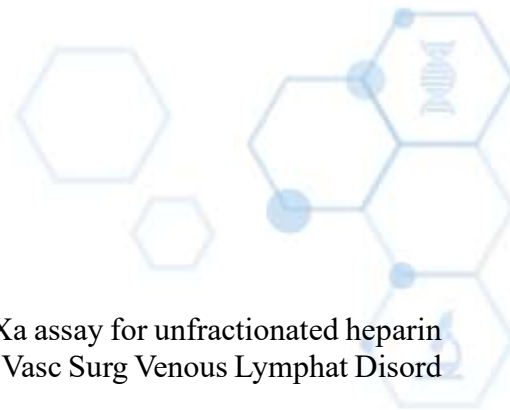
Mar 2015 – Feb 2017 Clinical Fellow,
Department of Laboratory Medicine and Genetics,
Samsung Medical Center, Seoul, Korea

Mar 2011 – Feb 2015 Residency,
Department of Laboratory Medicine and Genetics,
Samsung Medical Center, Seoul, Korea

• **Professional Organizations:**

- Member, The Korean Society for Laboratory Medicine
- Member, The Korean Society of Hematology
- Member, The Korean Society of Diagnostic Hematology
- Member, The Korean Society of Genetic & Molecular Diagnosis





• Main Scientific Publications:

1. Shin H, Koh EH, Lee GW, Song HN, Kim HY, Park S. Can an anti-Xa assay for unfractionated heparin be used to assess the presence of rivaroxaban in critical situations? *J Vasc Surg Venous Lymphat Disord* 2020.
2. Kim HY, Chung YN, Cho D. Extended Red Blood Cell Phenotype Matching Is Dependent on Ethnicity and Specificity of RBC Alloantibodies. *Ann Lab Med* 2020;40:190-2.
3. Shin H, Park S, Lee GW, Koh EH, Kim HY. Parvovirus B19 infection presenting with neutropenia and thrombocytopenia: Three case reports. *Medicine (Baltimore)* 2019;98:e16993.
4. Park CH, Yun JW, Kim HY, Lee KO, Kim SH, Kim HJ. Myelodysplastic Syndrome/Myeloproliferative Neoplasm with Ring Sideroblasts and Thrombocytosis with Cooccurrent SF3B1 and MPL Gene Mutations: A Case Report and Brief Review of the Literature. *Lab Med* 2019.
5. Kim HY, Park SW, Kim JH, Kang JH, Lee WS, Song HN. Romiplostim-related myelofibrosis in refractory primary immune thrombocytopenia: A Case report. *Medicine (Baltimore)* 2019;98:e15882.
6. Kim HY, Kim JH, Jung MH, Cho IA, Kim Y, Cho MC. Clinical Usefulness of Bioavailable Vitamin D and Impact of GC Genotyping on the Determination of Bioavailable Vitamin D in a Korean Population. *Int J Endocrinol* 2019;2019:9120467.
7. Kim HY, Cho EJ, Chun S, Kim KH, Cho D. Red Blood Cell Alloimmunization in Korean Patients With Myelodysplastic Syndrome and Liver Cirrhosis. *Ann Lab Med* 2019;39:218-22.
8. Choi JY, Kim HY, Kang MG, Shin JK, Lee WS, Song HN. Bilateral ovarian granulocytic sarcoma as the primary manifestation of acute myelogenous leukemia treated with allogenic stem cell transplantation: A case report. *Medicine (Baltimore)* 2019;98:e18390.
9. Kim HY, Lee KO, Park S, Jang JH, Jung CW, Kim SH, et al. Poor Prognostic Implication of ASXL1 Mutations in Korean Patients With Chronic Myelomonocytic Leukemia. *Ann Lab Med* 2018;38:495-502.
10. Kim HY, Kim JW, Kim SH, Chang MH, Kim HJ. A novel type of +2-base pair frameshift CALR mutation in a patient with myeloproliferative neoplasm. *Eur J Haematol* 2018.
11. Kim HY, Cho JH, Kim HJ, Chung HS, Kim SH, Lee KO, et al. Ethnicity-specific impact of HLA I/II genotypes on the risk of inhibitor development: data from Korean patients with severe hemophilia A. *Ann Hematol* 2018.
12. Shin SY, Kim HY, Kim HJ, Kim HG. Hb Heathrow [beta103(G5)Phe-->Leu], a First Report in an Asian Patient with Erythrocytosis. *Yonsei Med J* 2017;58:665-7.
13. Kim HY, Park S, Kim SH, Kim HJ. The first report of chronic myelogenous leukemia with e19a2 micro-BCR/ABL1 presenting with massive myelofibrosis. *Blood Cells Mol Dis* 2017;65:68-70.
14. Kim HY, Jang MA, Kim HJ, Kim SJ, Kim WS, Kim SH. Clinical impact of CD5 expression in Korean patients with diffuse large B-cell lymphoma. *Blood Res* 2017;52:193-9.
15. Chun S, Phan MT, Kim HY, Shin DJ, Seo JY, Kim KH, et al. The A312 Allele (c.280A>T) Is Responsible for the Weak A Phenotype. *Ann Clin Lab Sci* 2017;47:99-102.
16. Shin SY, Lee ST, Kim HY, Park CH, Kim HJ, Kim JW, et al. Detection of MYD88 L265P in patients with lymphoplasmacytic lymphoma/Waldenstrom macroglobulinemia and other B-cell non-Hodgkin lymphomas. *BLOOD RESEARCH* 2016;51:181-6.
17. Kim HY, Lee KO, Yoo KH, Kim SH, Oh D, Kim HJ. Congenital thrombotic thrombocytopenic purpura (Upshaw-Schulman syndrome) caused by novel ADAMTS13 mutations. *Br J Haematol* 2016;173:156-9.
18. Kim HY, Lee SH, Lee MN, Kim JW, Kim YH, Kim MJ, et al. Complete sequence-based screening of TPMT variants in the Korean population. *Pharmacogenet Genomics* 2015;25:143-6.