

Manuscript no. HAEMATOL/2011/045591 entitled “Disruption of the ASXL1 gene is frequent in primary, post-essential thrombocytosis and post-polycythemia vera myelofibrosis, but not essential thrombocytosis or polycythemia vera: analysis of molecular genetics and clinical phenotypes”

Authors: Brady L. Stein, Donna M. Williams, Christine O’Keefe, Ophelia Rogers, Roxann G. Ingersoll, Jerry L. Spivak, Amit Verma, Jarek P. Maciejewski, Michael A. McDevitt, Alison R. Moliterno

Information about the contributions of each person named as having participated in the study

1) Guarantor(s), i.e., person(s) who is (are) responsible for the integrity of the work as a whole:
• Alison R. Moliterno MD, Johns Hopkins University School of Medicine, amoliter@jhmi.edu

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2) Authors who participated in the conception of the study: Brady Stein, Donna Williams, Christine O’Keefe, Amit Verma, Jarek Maciejewski, Jerry Spivak, Michael McDevitt, and Alison Moliterno

3) Design & Methods. The following authors were responsible for specific investigations :
• Brady Stein, Jerry Spivak, Michael McDevitt and Alison Moliterno were responsible for designing the clinical database, analyzing mutation frequencies and performing the statistical analyses.
• Donna Williams, Ophelia Rogers, Christine O’Keefe, Roxann Ingersoll and Alison Moliterno were responsible for experimental design of genomic analyses including SNP-array karyotyping, JAK2V617F genotyping, direct sequencing and pyrosequencing

4) Results. The following authors were responsible for specific portions of the results, including figures and tables:
• Brady Stein and Alison Moliterno were responsible for Tables 1, 4 and 5
• Alison Moliterno and Donna Williams were responsible for Tables 2 and 3 and all the Figures
• Alison Moliterno, Ophelia Rogers, Donna Williams, Christine O’keefe and Roxann Ingersoll were responsible for interpretation of SNP-array karyotyping, genomic sequence variation and pyrosequencing results

5) Writing the manuscript. The following authors were responsible for writing the manuscript:
• Brady Stein, Donna Williams, Michael McDevitt and Alison Moliterno drafted and edited the manuscript
• Christine O’keefe, Jerry Spivak, Amit Verma, and Jarek Maciejewski edited the manuscript

6) Contributors Listed in Acknowledgments:

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