

haematologica *the hematology journal*

author contribution form

Title	In rare acute myeloid leukemia patients harboring both <i>RUNX1</i> and <i>NPM1</i> mutations, <i>RUNX1</i> mutations are unusual in structure and present in the germline
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Specific contributions of each author are listed on page 2 of this document.

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Contributions:

Clara D. Bloomfield conceived of the project, analyzed/interpreted data, and helped prepare the text.

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