

# haematologica *the hematology journal*

## author contribution form

<b>Title</b>	SF3B1 mutations are infrequently found in non-Myelodysplastic bone marrow failure syndromes and mast cell diseases but, if present, are associated with the ring sideroblast phenotype
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*Specific contributions of each author are listed on page 2 of this document.*

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*Please describe the contributions of each author, indicating who was responsible for each part of the study and the preparation of the manuscript (collection of data, experiments, data analysis, generation of figures, collection of images, interpretation of data, preparation of the text, etc.)*

Valeria Visconte designed the study, performed experiments, analyzed and interpreted the data, and wrote the manuscript.

Ali Tabaroki collected the data, performed experiments, and approved the manuscript.

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