

**Manuscript no. HAEMATOL/2011/054361 entitled “Germ-line GATA2 p.THR354MET mutation in familial myelodysplastic syndrome with acquired monosomy 7 and ASXL1 mutation demonstrating rapid onset and poor survival”**

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

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2) **Authors who participated in the conception of the study:** Csaba Bödör, Matthew Smith, Claude Preudhomme, Jude Fitzgibbon, Carolyn Owen

3) **Design & Methods.** The following authors were responsible for specific investigations:

- Csaba Bödör, Aline Renneville, Aurélie Charazac, Pascaline Étancelin, Sameena Iqbal and Karolina Kramarzová were responsible for collecting the samples and performing the molecular analyses.
- Matthew Smith, Jamie Cavenagh, András Matolcsy, Biju Krishnan and Carolyn Owen were responsible for collecting clinical data.

4) **Results.** The following authors were responsible for specific portions of the results, including figures and tables:

- Csaba Bödör and Carolyn Owen were responsible for Figures 1 and 2.
- Csaba Bödör, Aline Renneville, Aurélie Charazac and Karolina Kramarzová were responsible for the mutation data.
- Matthew Smith, Jamie Cavenagh, Sameena Iqbal and Carolyn Owen were responsible for analyzing the clinical data.

5) **Writing the manuscript.** The following authors were responsible for writing the manuscript:

- All authors were responsible for writing or critical revision of the article.

**6) Contributors Listed in Acknowledgments:**

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