

AUTHOR CONTRIBUTION FORM

Title	A Phase I trial of ribavirin and low-dose cytarabine for the treatment of relapsed and refractory acute myeloid leukemia with elevated eIF4E
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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, interpretation of data, preparation of the text, etc.):

SA was the principal investigator of the clinical trial. SA also contributed to the conception and design of the study as well as the drafting of the manuscript, participated in the collection and assembly of the data, recruited patients, analyzed and interpreted data. BC-K and HAZ designed, conducted, and analyzed experiments, as well as interpreted the data. EC coordinated all activities from multiple sites, contributed to the study design, drafting of the manuscript, collection, assembly, analysis and interpretation of the data. JB and SC contributed to patient recruitment. CL coordinated all activities from multiple sites and contributed to data collection and assembly. CJL contributed to data collection and assembly. WHM contributed to study design and analyzed data. KLBB was the principle investigator of the correlative studies. KLBB came up with the concept of targeting eIF4E with ribavirin in cancer, contributed to the conception and design of the study, data analysis, data assembly, interpretation of data and drafting of the manuscript. All authors revised the manuscript critically.