

Haematologica
HAEMATOL/2018/194670
Version 3

Contribution of alternative complement pathway to delayed hemolytic
transfusion reaction in sickle cell disease

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Disclosures: none

Contributions: S.C. collected and analyzed the data and wrote the manuscript. C.M.B.,
C.L.D., and M.O.Q. collected and analyzed the data and reviewed the manuscript. C.H.J.,
R.M.F., and S.R.S. analyzed the data and provided critical revisions to the manuscript.