Haematologica HAEMATOL/2016/160192 Version 4

Exome sequencing identifies recurrent BCOR gene alterations and the absence of KLF2, TNFAIP3 and MYD88 mutations in splenic diffuse red pulp small B-cell lymphoma

Laurent Jallades, Lucile Baseggio, Pierre Sujobert, Sarah Huet, Kaddour Chabane, Evelyne Callet-Bauchu, Aurélie Verney, Sandrine Hayette, Jean-Pierre Desvignes, David Salgado, Nicolas Levy, Christophe Béroud, Pascale Felman, Françoise Berger, Jean-Pierre Magaud, Laurent Genestier, Gilles Salles, and Alexandra Traverse-Glehen

Disclosures: All the authors declare no conflicts of interests.

Contributions: LJ performed the research, interpreted the data and wrote the paper; AV and KC performed the experiments; PS, SHu, SHa, JPD, CB, DS, NL, LG interpreted the data; clinical cases were selected and extensively reviewed by expert pathologists (ATG, FB), cytologists and immunologists (LJ, LB, PF), and cytogeneticists (ECB, JPM); LJ, ATG and GS designed the research study. LJ, LB, ATG, and GS interpreted the data and wrote the paper.