Time for newer approach in age-old AIHA

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Foremost, I would like to congratulate and thank authors of paper published in Lancet Regional Health, South East Asia to raise concerns on management of autoimmune hemolytic anemia (AIHA), which is usually considered as a benign haematological disorder.1 It has been rightly emphasized that there are many challenges pertaining to even diagnose correct type of AIHA. I support the fact that laboratories in India are not well equipped to perform cold agglutinin titers, Donath-Landsteiner test, testing for drug induced AIHA even in tertiary care settings. In terms of treatment modalities, rituximab and complement inhibitors have been highlighted as major darts in armamentarium depicting complete response rate with single agent rituximab as low as 5%, extending to 53% when combined with bortezomib and bendamustine. Also, rituximab requires multiple doses before response is apparent. On the other hand, complement inhibitors are out of reach for majority of Indian patients till date. I aim to bring attention to anti-CD38 monoclonal antibody 'Daratumumab' as a measure to reduce autoantibodies at a faster rate. In my clinical experience, Daratumumab has provided 100% remission even at first dose, particularly when cytopenia is severe and multilineage like in Evan's syndrome. There is substantial literature available which supports its effectiveness with very little toxicity. Hence, it should be considered in steroid refractory patients as a second line of treatment. I firmly believe that Daratumumab is worthy of attention to be a part of AIHA treatment algorithm, as its usefulness has been widely experienced and acknowledged.

Contributors

PM-conceptualised, drafted, wrote, revised and edited the manuscript.

Declaration of interests

There is no conflict of interest to declare.

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