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# **Clinical Updates on Haemophilia**

Guest Editor:

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Deadline for manuscript submissions: **31 January 2025** 

### Message from the Guest Editor

Dear Colleagues,

Over the last decade, the number of products used to treat a person with haemophilia (PwH) has grown dramatically. Novel therapies offer both improved protection from bleeds and more comfort of administration. Extended halflife products allow for an increase in factor levels, even reducing the number of infusions, especially in haemophilia B. The requirement of less infusions licenses the start of regular prophylaxis even in moderate haemophilia patients. In severe haemophilia A patients, subcutaneous non-replacement therapy permits the start of prophylaxis very early in children's lives, as intravenous administration is no longer a problem. Lastly, gene therapy is now available, and PwH who undergo gene therapy may see their factor levels increase, sometimes even to normal levels.



mdpi.com/si/201990







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