

Amy Shapiro

List of Publications by Year in descending order

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Version: 2024-02-01

22
papers

1,851
citations

516215

16
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676716

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all docs

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docs citations

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times ranked

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citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | The effect of emicizumab prophylaxis on long-term, self-reported physical health in persons with haemophilia A without factor VIII inhibitors in the HAVEN 3 and HAVEN 4 studies. <i>Haemophilia</i> , 2021, 27, 854-865. | 1.0 | 21 |
| 2 | Final results of the PUPs B-LONG study: evaluating safety and efficacy of rFIXFc in previously untreated patients with hemophilia B. <i>Blood Advances</i> , 2021, 5, 2732-2739. | 2.5 | 11 |
| 3 | The use of prophylaxis in the treatment of rare bleeding disorders. <i>Thrombosis Research</i> , 2020, 196, 590-602. | 0.8 | 23 |
| 4 | Real-world data demonstrate improved bleed control and extended dosing intervals for patients with haemophilia B after switching to recombinant factor IX Fc fusion protein (rFIXFc) for up to 5 years. <i>Haemophilia</i> , 2020, 26, 975-983. | 1.0 | 12 |
| 5 | Long-term safety and sustained efficacy for up to 5 years of treatment with recombinant factor IX Fc fusion protein in subjects with haemophilia B: Results from the BOND extension study. <i>Haemophilia</i> , 2020, 26, e262-e271. | 1.0 | 28 |
| 6 | Young adult outcomes of childhood prophylaxis for severe hemophilia A: results of the Joint Outcome Continuation Study. <i>Blood Advances</i> , 2020, 4, 2451-2459. | 2.5 | 67 |
| 7 | Hemophilia A with inhibitor: Immune tolerance induction (ITI) in the mirror of time. <i>Transfusion and Apheresis Science</i> , 2019, 58, 578-589. | 0.5 | 17 |
| 8 | Using pharmacokinetics for tailoring prophylaxis in people with hemophilia switching between clotting factor products: A scoping review. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2019, 3, 528-541. | 1.0 | 18 |
| 9 | Efficacy, safety, and pharmacokinetics of emicizumab prophylaxis given every 4 weeks in people with haemophilia A (HAVEN 4): a multicentre, open-label, non-randomised phase 3 study. <i>Lancet Haematology</i> , 2019, 6, e295-e305. | 2.2 | 252 |
| 10 | BIVV001: The First Investigational Factor VIII Therapy to Break Through the VWF Ceiling in Hemophilia A, with Potential for Extended Protection for One Week or Longer. <i>Blood</i> , 2018, 132, 636-636. | 0.6 | 11 |
| 11 | Plasma-derived human factor X concentrate for on-demand and perioperative treatment in factor X-deficient patients: pharmacology, pharmacokinetics, efficacy, and safety. <i>Expert Opinion on Drug Metabolism and Toxicology</i> , 2017, 13, 97-104. | 1.5 | 25 |
| 12 | Safety and efficacy of recombinant factor VIIa by pediatric age cohort: reassessment of compassionate use and trial data supporting US label. <i>Pediatric Blood and Cancer</i> , 2016, 63, 1822-1828. | 0.8 | 8 |
| 13 | Switching to recombinant factor IX Fc fusion protein prophylaxis results in fewer infusions, decreased factor IX consumption and lower bleeding rates. <i>British Journal of Haematology</i> , 2015, 168, 113-123. | 1.2 | 31 |
| 14 | Phase 3 study of recombinant factor VIII Fc fusion protein in severe hemophilia A. <i>Blood</i> , 2014, 123, 317-325. | 0.6 | 403 |
| 15 | Development of long-acting recombinant FVIII and FIX Fc fusion proteins for the management of hemophilia. <i>Expert Opinion on Biological Therapy</i> , 2013, 13, 1287-1297. | 1.4 | 33 |
| 16 | Association Of Bleeding Tendency With Time Under Target FIX Activity Levels In Severe Hemophilia B Patients Treated With Recombinant Factor IX Fc Fusion Protein. <i>Blood</i> , 2013, 122, 2349-2349. | 0.6 | 9 |
| 17 | Integrated analysis of safety and efficacy of a plasma- and albumin-free recombinant factor VIII (rAHF-PFM) from six clinical studies in patients with hemophilia A. <i>Expert Opinion on Biological Therapy</i> , 2009, 9, 273-283. | 1.4 | 31 |
| 18 | Surgical evaluation of a recombinant factor VIII prepared using a plasma/albumin-free method: Efficacy and safety of Advate in previously treated patients. <i>Thrombosis and Haemostasis</i> , 2008, 100, 217-223. | 1.8 | 60 |

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|----|---|-----|-----------|
| 19 | Home Treatment of Mild to Moderate Bleeding Episodes Using Recombinant Factor VIIa (Novoseven) in Haemophiliacs with Inhibitors. <i>Thrombosis and Haemostasis</i> , 1998, 80, 912-918. | 1.8 | 350 |
| 20 | Prospective, Randomised Trial of Two Doses of rFVIIa (NovoSeven) in Haemophilia Patients with Inhibitors Undergoing Surgery. <i>Thrombosis and Haemostasis</i> , 1998, 80, 773-778. | 1.8 | 365 |
| 21 | The pattern of spontaneous germ-line mutation: relative rates of mutation at or near CpG dinucleotides in the factor IX gene. <i>Human Genetics</i> , 1993, 91, 496-503. | 1.8 | 42 |
| 22 | A past mutation at Isoleucine397 is now a common cause of moderate/mild haemophilia B. <i>British Journal of Haematology</i> , 1990, 75, 212-216. | 1.2 | 34 |