












ORIGINAL ARTICLE

F9 missense mutations impairing factor IX activation are associated with pleiotropic plasma phenotypes

Alessio Branchini¹  | Massimo Morfini²  | Barbara Lunghi¹  | Donata Belvini³ | Paolo Radossi⁴ | Loredana Bury⁵  | Maria Luisa Serino⁶ | Paola Giordano⁷  | Dorina Cultrera⁸ | Angelo Claudio Molinari⁹  | Mariasanta Napolitano¹⁰  | Elisabetta Bigagli¹¹  | Giancarlo Castaman¹²  | Mirko Pinotti¹  | Francesco Bernardi¹  | The GePKHIS Study Group of AICE

¹Department of Life Sciences and Biotechnology and LTTA Centre, University of Ferrara, Ferrara, Italy

²Italian Association of Haemophilia Centers (AICE), Milan, Italy

³Transfusion Service, Haemophilia Centre and Haematology, Castelfranco Veneto Hospital, Castelfranco Veneto, Italy

⁴Oncohematology-Oncologic Institute of Veneto, Castelfranco Veneto Hospital, Castelfranco Veneto, Italy

⁵Department of Medicine and Surgery, University of Perugia, Perugia, Italy

⁶Haemostasis and Thrombosis Centre, University Hospital of Ferrara, Ferrara, Italy

⁷Paediatric Section, Department of Biomedicine and Human Oncology, A. Moro" University, Bari, Italy

⁸Haemophilia Regional Reference Center, Vittorio Emanuele" University Hospital, Catania, Italy

⁹Thrombosis and Haemostasis Unit, Gaslini Hospital, Genova, Italy

¹⁰Haematology Unit, Thrombosis and Haemostasis Reference Regional Center and PROMISE Department, University of Palermo, Palermo, Italy

¹¹Department of Neuroscience, Psychology, Drug Research and Child Health (NEUROFARBA), Section of Pharmacology and Toxicology, University of Florence, Florence, Italy

¹²Department of Oncology, Center for Bleeding Disorders, Careggi University Hospital, Firenze, Italy

Correspondence

Alessio Branchini and Francesco Bernardi,
Department of Life Sciences and
Biotechnology, University of Ferrara, Via
Fossato di Mortara 74, 44121 Ferrara,
Italy.

Emails: brnlss@unife.it (AB);

ber@unife.it (FB)

Funding information

Pfizer, Grant/Award Number: WI225911

Abstract

Background: Circulating dysfunctional factor IX (FIX) might modulate distribution of infused FIX in hemophilia B (HB) patients. Recurrent substitutions at FIX activation sites (R191-R226, >300 patients) are associated with variable FIX activity and antigen (FIXag) levels.

Objectives: To investigate the (1) expression of a complete panel of missense mutations at FIX activation sites and (2) contribution of F9 genotypes on the FIX pharmacokinetics (PK).

Methods: We checked FIX activity and antigen and activity assays in plasma and after recombinant expression of FIX variants and performed an analysis of infused FIX PK

Manuscript handled by: James Morrissey

Final decision: James Morrissey, 06 October 2021

Branchini, Morfini, and Lunghi equally contributed to this study.

The members of GePKHIS Study group of AICE are listed in the Appendix.

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

© 2021 The Authors. *Journal of Thrombosis and Haemostasis* published by Wiley Periodicals LLC on behalf of International Society on Thrombosis and Haemostasis

parameters in patients ($n = 30$), mostly enrolled in the *F9* Genotype and PK HB Italian Study (GePKHIS; EudraCT ID2017-003902-42).

Results: The variable FIXag amounts and good relation between biosynthesis and activity of multiple R191 variants results in graded moderate-to-mild severity of the R191C>L>P>H substitutions. Recombinant expression may predict the absence in the HB mutation database of the benign R191Q/W/K and R226K substitutions. Equivalent changes at R191/R226 produced higher FIXag levels for R226Q/W/P substitutions, as also observed in p.R226W female carrier plasma.

Pharmacokinetics analysis in patients suggested that infused FIX Alpha distribution and Beta elimination phases positively correlated with endogenous FIXag levels. Mean residence time was particularly prolonged (79.4 h, 95% confidence interval 44.3–114.5) in patients ($n = 7$) with the R191/R226 substitutions, which in regression analysis were independent predictors (β coefficient 0.699, $P = .004$) of Beta half-life, potentially prolonged by the increasing over time ratio between endogenous and infused FIX.

Conclusions: FIX activity and antigen levels and specific features of the dysfunctional R191/R226 variants may exert pleiotropic effects both on HB patients' phenotypes and substitutive treatment.

KEYWORDS

factor IX activation, hemophilia B, pharmacogenetics, pharmacokinetics, recombinant proteins

1 | INTRODUCTION

The activation of coagulation factor IX (FIX) has a key role in the coagulation cascade, either via activated factor VII (FVIIa) in complex with tissue factor (TF) or via activated factor XI (FXIa).¹ Full activation is achieved after cleavage at two highly conserved activation sites,^{2,3} which results in the release of the activation peptide.⁴⁻⁷ More than 300 hemophilia B (HB) patients with recurrent *F9* mutations affecting the R191 or R226 residues, encoded by CpG-containing codons,⁸ have been reported (EAHAD FIX Variant Database, <https://f9-db.eahad.org/>).⁹ Seminal studies have indicated that single cleavage at R191 does not produce catalytic activity but converts FIX zymogen into a factor VIII (FVIII)-binding enzyme,^{10,11} whereas single cleavage at R226 develops catalytic activity but results in suboptimal binding to the FVIII light chain.^{11,12} Accordingly, missense mutations at these positions are associated in HB patients with moderate/mild (R191)¹³⁻²¹ or severe (R226, HB_m mutation subclass)^{14,22-28} FIX deficiency. Failure in the production of activated FIX (FIXa)^{14,24,29} has been modelled in HB mice (R226W).³⁰

Based on this information, we focused on *F9* mutations affecting the FIX activation sites by systematic recombinant expression of natural and designed variants to interpret HB phenotypes, both residual activity and antigen levels, associated to virtually all amino acid substitutions.

Previous results suggested that activation peptide sequences act as plasma retention signals, as observed for human FVII added

Essentials

- The factor IX (FIX) cross-reactive material (CRM) might modulate infused FIX distribution.
- The contribution of *F9* genotypes on infused FIX pharmacokinetics was investigated.
- FIX activation site (R191/226) variants (hemophilia B patients $n > 300$) express medium/high cross-reactive material levels.
- FIX activation site substitutions may prolong infused FIX Alpha and Beta half-lives and mean residence time.

with the FIX activation peptide, which prolonged 4.3-fold its half-life when tested after infusion in mice.^{31,32}

Concerning the distribution of FIX infused in HB patients, it has been hypothesized that the presence of cross-reactive material (CRM⁺), namely the circulating FIX antigen (FIXag) with reduced or null activity, might partially improve the *in vivo* recovery of FIX infused in HB patients.³³ Differently, in HB mice, the presence of dysfunctional endogenous FIX caused by the human R379Q variant³⁴ may decrease replacement efficacy.³⁵ Based on these observations, it has been proposed that prophylaxis regimens in HB patients should consider CRM status.³⁶ Because observations in patients and animal models may be explained by the dynamic equilibrium between plasma FIX and the extravascular protein bound to the subendothelial basement

membrane,³⁷⁻⁴⁰ a two-compartment (2CP) pharmacokinetic (PK) model best describes the distribution of FIX infused in HB. We report the PK analysis of FIX infused in severe/moderate HB patients, enrolled in the frame of the *F9* Genotype and PK Haemophilia B Italian Study (GePKHIS), to investigate the relation between distribution of infused FIX and circulating endogenous FIX variants, among which there are several substitutions at the FIX activation sites.

2 | MATERIALS AND METHODS

2.1 | Patients and approval

Plasma samples from HB patients ($n = 30$), mostly enrolled in the GePKHIS study, and in addition three related female carriers of the p.R226W variant, were collected and processed at hemophilia centers belonging to the Italian Association of Haemophilia Centres (AICE). The study was approved by the coordinator center (Ferrara, code ACTB02BD04), as well as by each local ethics committee, and was registered in the EudraCT database (ID 2017-003902-42). The study was carried out in accordance with the Declaration of Helsinki, and written informed consent was obtained.

2.2 | Study design

The protocol was approved for severe/moderate HB patients (Table S1 in supporting information) undergoing on-demand or prophylactic treatment with nonacog alfa (Pfizer).

Inclusion criteria to participate in the study were: severe or moderately severe HB (FIX activity ≤ 3 IU/dL); no bleeds in the last week before assessment of PK; on-demand or on-prophylaxis treatment with nonacog alfa for >150 exposure days.

Exclusion criteria were: any kind of bleeding in the last week before the PK, severe hepatic disease, ongoing human immunodeficiency virus treatment (highly active antiretroviral treatment), history of anti-FIX inhibitors, previous treatments with extended half-life FIX concentrates. The PK study was not conducted in the HB patient with the p.R191H variant, associated with a mild HB phenotype (FIX coagulant activity $>5\%$; see Table S1). For this patient, only the basal activity and antigen levels were determined.

2.3 | Nomenclature

All amino acids are numbered according to the Human Genome Variation Society (HGVS) nomenclature.⁴¹

2.4 | *F9* genotypes of patients

The *F9* genotypes of HB patients undergoing PK (Table S1) were characterized as described.²⁰ Causative mutations were classified as

missense ($n = 16$, patients $n = 21$), nonsense ($n = 4$), splicing ($n = 2$), and deletion ($n = 2$) variants.

2.5 | Expression of recombinant variants

Recombinant FIX (rFIX) variants were created by site-directed mutagenesis of *F9* cDNA (reference sequences: NM_000133.4, NP_000124.1) cloned in the pCDNA3 vector.⁴² Oligonucleotides are listed in Table S2 in supporting information. All plasmids have been validated by sequencing. Nonsense variants have been produced in previous studies.^{43,44}

Expression studies were carried out through transient transfection of human embryonic kidney 293 (HEK293) cells in the presence of $5 \mu\text{g/mL}$ vitamin K (konakion, 10 mg/mL), essentially as described.⁴² Briefly, cells were seeded in 12-well culture plates and transfected in serum-free medium (Opti-MEM, Gibco, Life Technologies) with the Lipofectamine 2000 reagent (Life Technologies) with a ratio DNA (μg):Lipofectamine (μL) of 1:1 ($2 \mu\text{g}$ DNA: $2 \mu\text{L}$ Lipofectamine) and media were collected 48 h post-transfection.

2.6 | Evaluation of protein and activity of rFIX variants

FIX activity and antigen levels were evaluated by polyclonal anti-human FIX ELISA (FIX-EIA, Affinity Biologicals)^{43,44} to minimize the effects of mutations on detection of FIX epitopes. Known concentrations of purified rFIX were used as reference.

Protein forms derived from FXIa-dependent activation⁴⁵ of rFIX variants were evaluated through western blotting analysis with polyclonal goat anti-human FIX (APGAFIX; Affinity Biologicals) and anti-goat horseradish peroxidase-conjugated (A50-101P; Bethyl Laboratories) antibodies. Blotting images were acquired on the ChemiDoc instrument and analyzed by the Image Laboratory Software version 4.0 (Bio-Rad).

Activity of rFIX variants in media was evaluated through a commercially available chromogenic assay exploiting an optimized FXIa concentration as per manufacturer instructions (Biophen; Aniaara Diagnostica).⁴³ Reaction aliquots were collected for western blotting analysis. Serial dilutions of wild-type rFIX (rFIX-WT, mean concentration $485 \pm 51.5 \text{ ng/mL}$), expressed in the same experimental conditions in parallel to rFIX variants, were used as reference. The activity/antigen ratio was calculated.

2.7 | Evaluation of FIX protein and activity in plasma

Serial dilutions of pooled normal plasma (PNP; Hyphen BioMed) were used as reference for analysis through polyclonal anti-human FIX ELISA (Affinity Biologicals)^{43,44} of FIXag levels in plasma from

HB patients. Activity of infused/baseline FIX was measured by one-stage clotting method.⁴⁶

2.8 | PK analysis

Each patient received a single dose (40 ± 5 IU/kg) of nona-cog alfa. Plasma samples were collected before infusion and at 1–3–9–24–48–72 h post-infusion.

PK parameters were obtained through the 2CP model analysis (WinNonlin 7.0),^{46–48} with the best fitting evaluated through the sum of squared residuals (SSR)⁴⁹ and the coefficient of correlation between observed and predicted FIX concentration/time. The FIX/time concentrations were corrected according to the Björkman formula⁵⁰ or by basal level subtraction, as appropriate.

Specific primary (K 1–2, K 2–1, and V1) and secondary (Alpha HL, Beta HL, Clearance, CLD2, Cmax, mean residence time [MRT], V2, Vss and area under the curve [AUC]) PK parameters, including *in vivo* recovery (U/dL per IU FIX concentrate/kg body weight infused) were evaluated.^{46,51,52}

2.9 | Statistical analysis

Statistical analyses were performed using IBM® SPSS® Statistics (version 23.0; IBM Corp.). Variables were reported as means with 95% confidence interval (CI). For skewed variables, the analysis was conducted with logarithmically transformed values. Two-tailed Pearson correlations were performed to determine the association between PK parameters and FIXag (%). F9 mutation type-related differences in PK parameters were analyzed by *t*-test. In linear regression analysis, the contribution to PK parameters of FIXag levels and F9 mutation types was evaluated.

3 | RESULTS

3.1 | R191/R226 substitutions, plasma, and recombinant phenotypes

To compare expression levels of protein variants with amino acid substitutions at FIX activation sites (R191 and R226), we evaluated FIXag and activity levels obtained from (1) analysis of HB patients' plasma, collected in the frame of the GePKHIS study; (2) inspection of the international database (<https://f9-db.eahad.org/>); and (3) systematic expression of rFIX variants harboring natural or designed changes.

In HB patients (Figure 1A), the R191H/C substitutions result in reduced plasma FIX activity (Figure 2A, left panel) and normal (R191H) or moderately reduced (R191C) FIXag (Figure 2A, right panel) levels. The potential effects of the unpaired cysteine in the 191C FIX were explored by western blotting analysis of plasma and conditioned medium performed in reducing/non-reducing

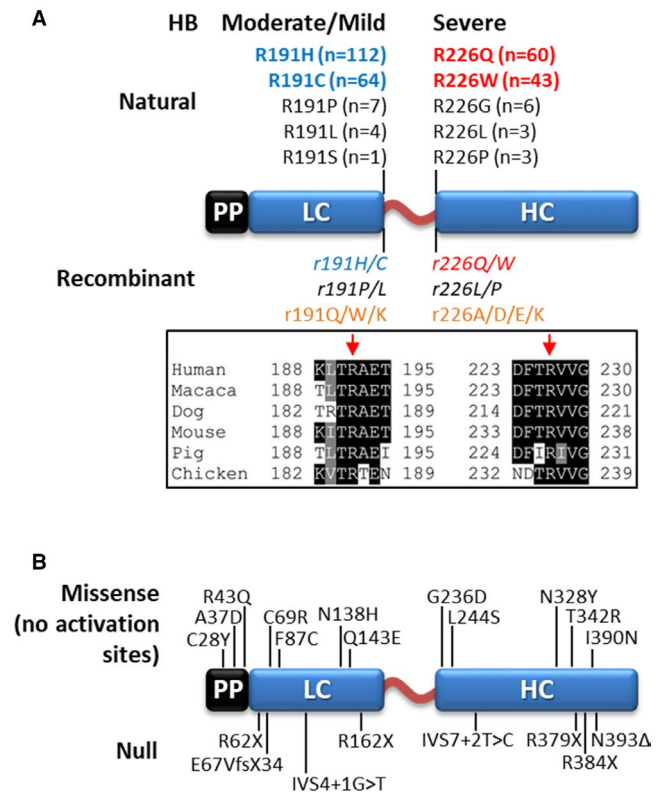


FIGURE 1 Natural and designed amino acid substitutions at factor IX (FIX) activation sites. A, Schematic representation of human coagulation FIX showing the pre- and pro-peptides (PP), and light and heavy chains (LC and HC, respectively). Hemophilia B (natural) variants affecting the R191 and R226 sites are reported above the FIX scheme (n, patient number in the EAHAD FIX Variant Database <https://f9-db.eahad.org/>),⁹ with the prevalent HB phenotypes associated with mutations on top. Blue and red variants, patients also available in the GePKHIS study. Designed recombinant variants are indicated below the FIX scheme. Color code: blue, red, and black, natural variants; orange, designed variants. The alignment and conservation of residues in the two activation sites (red arrows) is also shown. B, FIX variants available in the GePKHIS study. Missense (with amino acid changes not involving activation sites) and null (nonsense, splicing, and deletion) variants are reported above and below the FIX scheme, respectively

conditions. Data did not support the presence of high molecular weight molecules produced by disulfide bond of the 191C variant, either circulating in plasma or after secretion in medium (Figure S1 in supporting information). The devoid-of-function R226Q/W variants showed normal or increased antigen levels (Figure 2A, right panel). The increased plasma levels associated with the R226 substitutions is supported by data in HB patients and recombinant expression. Among the 60 patients affected by the R226Q and reported in the database, eight have been characterized for the FIXag, with mean levels >100%. Patients (n = 43) affected by the R226W change display (10 patients evaluated) variable FIXag levels (range 130–50%, mean 84%). Among the GePKHIS patients bearing these substitutions, FIXag was >100%.

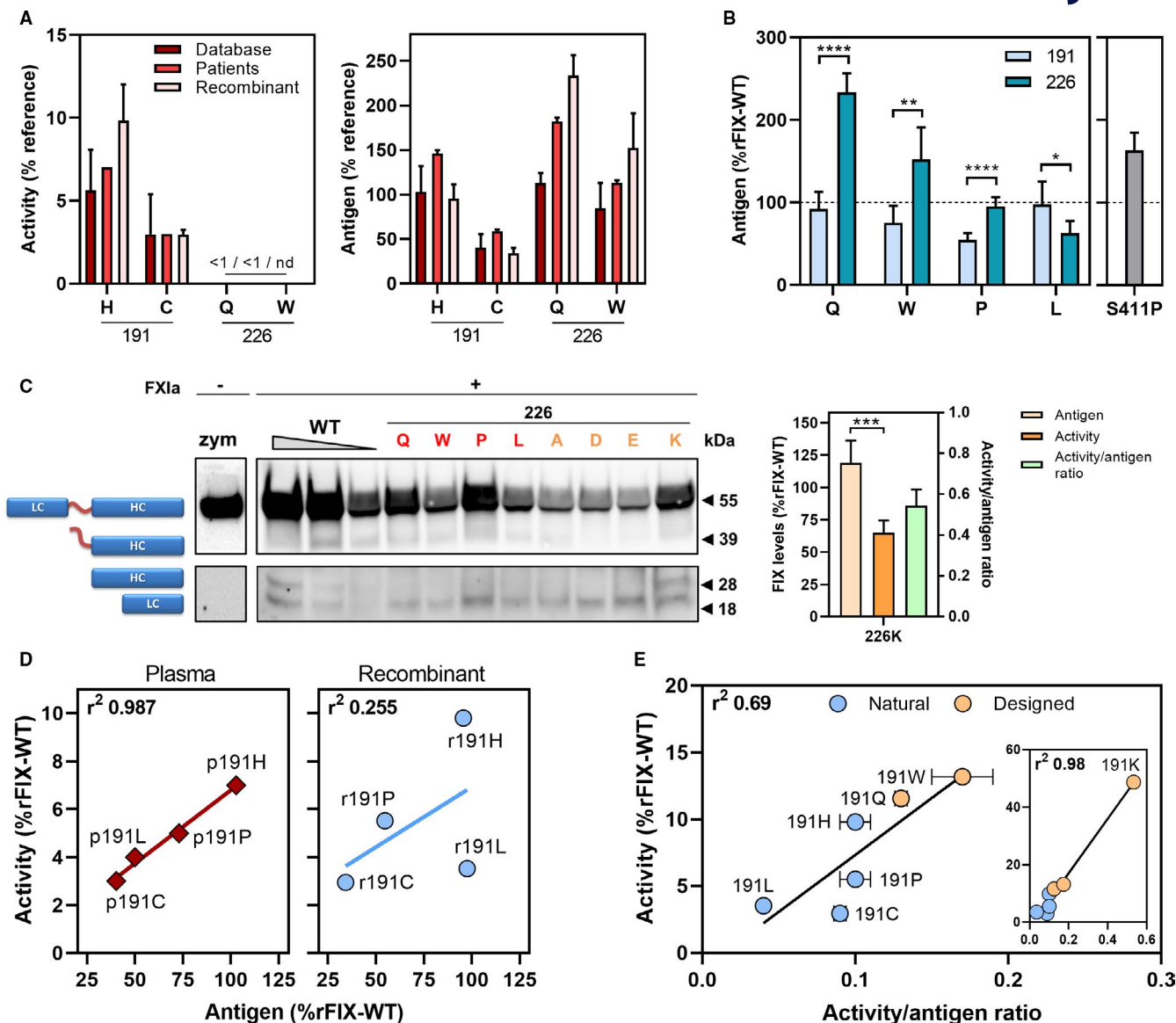


FIGURE 2 Characterization of factor IX (FIX) variants at the activation sites. **A**, Activity (left panel) and antigen (right panel) levels of the 191 and 226 variants estimated in GePKHIS patients, reported in the international EAHAD FIX variant database (<https://f9-db.eahad.org/>)⁹ or upon recombinant expression. Reference for plasma or recombinant values were pooled normal plasma or recombinant FIX wild type (rFIX-WT), as appropriate. **B**, Comparative expression of equivalent amino acid substitutions at 191 and 226 positions, after transient expression in HEK293 cells. The S411P variant (chymotrypsin numbering S195) was expressed as an additional control for inactive FIX with high antigen values. **C**, Left panel. Western blotting analysis of the activation profile of natural (red) and designed (orange) recombinant 226 variants in the presence (+) or absence (-) of FXIa. Protein forms, resulting from FIX activation, as well as relative molecular weights (kDa), are indicated on left and right of the blot, respectively. Serial dilutions of rFIX-WT were loaded as control and reference. rFIX variants with similar FIX antigen levels (226P/L/A/D/E/K, range 70%–100%) were tested after equal dilution (1:10) in phosphate buffered saline (PBS), while those with antigen levels exceeding wild-type rFIX (226Q/W), prior to being diluted in PBS, were first normalized to ~100% by dilution in medium from untransfected cells. Zym, zymogen FIX; FXIa, activated factor XI. Right panel. FIX antigen and activity levels, as well as activity/antigen ratio, of the designed 226K variant. **D**, Comparative analysis of activity and antigen levels of FIX variants bearing natural missense changes found in hemophilia B (HB) patients (p, red squares; left panel) and after recombinant expression (r, blue circles; right panel). **E**, Relation between mean activity levels and activity/antigen ratio of FIX variants bearing natural (blue circles) or designed (orange circles) amino acid substitutions. Inset, Analysis of the relation with the 191K variant included. Results, indicated as % of reference, are reported as mean \pm standard error of the mean (panels A, patients bar, and E) or mean \pm standard deviation of $n = 6$ replicates (panels A, recombinant bar, B and C). * $P < .05$; ** $P < .01$; *** $P < .001$; **** $P < .0001$

In expression studies, equivalent substitutions (R191/R226 to Q/W/P/L; Figure 2B) caused significantly different effects at position R191 and R226, and were compatible with normal or increased

secretion in media of the 226 variants compared to near normal or reduced levels of the 191 variants. As an additional control for catalytically inert serine proteases we also expressed the S411P variant

(chymotrypsin numbering S195), which also gave rise to high FIXag values.

Mutations affecting residues adjacent to R191 (A192P) and to R226 (V227F, V228L/F/A/G, Figure S2 in supporting information) are responsible for noticeable FIXag levels (70–130%), and for residual activity (2–20%) of most variants.^{17,27,53–55} Inspection of the exome sequencing database gnomAD (<https://gnomad.broadinstitute.org/>) revealed the presence of 12 subjects carrying the conservative V227I change (rs137852242), suggesting a rare polymorphism (Figure S2).

In expression studies we also designed and expressed rFIX control variants bearing a “classic” alanine change (R226A), or amino acid changes conferring negative (R226D/E) or positive (R226K) charge. Comparison between the natural (Q/W/P/L) and the designed (A/D/E, Figure S3 in supporting information) R226 variants in FXIa-dependent activation experiments (Figure 2C, left panel), failed to detect the FIX form corresponding to the heavy chain upon loss of the activation peptide. Differently, the designed R226K variant, as expected for the conservative R-to-K amino acid change, displayed an activation pattern similar to that of rFIX-WT (Figure 2C, left panel), and reached an activity and activity/antigen ratio above 50% and 0.5, respectively (Figure 2C, right panel).

To provide insights on the activity features of the R191C/L/P/H variants, the relation between FIXag and activity was investigated by comparing findings in plasma and recombinant systems (Figure 2D), which showed a very good relation only for plasma values ($r^2 = 0.987$). The normal FIXag levels for the r191L correspond to FIXag reduced to half in plasma, which suggests a FIX molecule with increased clearance *in vivo*. The relation between activity and activity/antigen ratio, which magnifies the functional properties of the multiple natural and designed R191 variants, was very good either excluding ($r^2 = 0.69$; Figure 2E) or including ($r^2 = 0.98$; Figure 2E, inset) the R191K variant, characterized by the highest activity and antigen values. The slope clearly indicated that the clinically relevant FIX deficiencies produced by the R191C/L substitutions (Figure 1A) are based on different functional properties. Conversely, the mild phenotype associated with the R191H deficiency is based on high secretion levels but modest catalytic properties.

Overall, comparison of *in vivo* and recombinant results of substitutions at activation sites provided a gradient of residual activity (R191) and antigen (both R191 and R226) values. The recombinant expression permits us to extrapolate information that is not available in patients but helps interpret findings in patients.

3.2 | Plasma FIX activity and antigen levels in female carriers of the p.R226W variant

We investigated plasma FIX activity and antigen levels in related female carriers^{56,57} ($n = 3$) of the p.R226W variant (Figure 3), to explore its potential influence on the proportion of the WT FIX

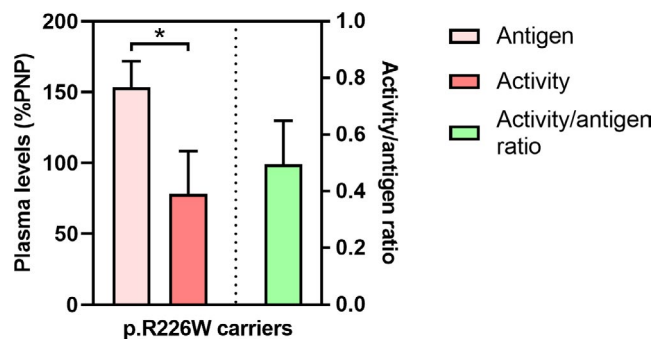


FIGURE 3 Analysis of female carriers of the p.R226W variant. Plasma factor IX (FIX) antigen and activity levels, as well as activity/antigen ratio, estimated in female carriers ($n = 3$) of the p.R226W variant. Results are reported as mean \pm standard error of the mean. * $P < .05$

and the inactive FIX variant circulating in plasma. The mean FIXag levels were higher ($153.4 \pm 18.6\%$) than the mean FIX activity ($78.0 \pm 30.3\%$), which resulted in an activity/antigen ratio decreased to half (0.50 ± 0.15 , Figure 3), suggesting similarly increased amounts of both FIX molecules.

3.3 | Correlation between FIX antigen levels and FIX PK parameters in HB patients

In the frame of the GePKHIS study, the decay of infused FIX was determined in 29 HB patients, characterized for *F9* genotypes. FIX variants corresponding to *F9* mutations in HB patients (Figure 1B and Table S1) were recombinantly expressed, which confirmed their causative role and supported the PK analysis. The characteristic biphasic decay curves and the SSR values are reported in Figure S4 in supporting information, and the distribution of 2CP PK parameter values in Table 1.

The FIXag levels were directly correlated (Table 1) with Alpha HL and Beta HL, and inversely with CLD2, Cmax, and with K 1–2 and K 2–1, the latter two primary parameters modelling the flow between plasma and extravascular compartments (Figure S4). The correlation between FIXag levels and *in vivo* recovery, expressed as the ratio between maximum FIX activity and the FIX dose, was not significant ($r = -0.289$, $P = .128$). On the other hand, the *in vivo* recovery has been reported to be quite variable even in the same patient on different occasions.⁵²

3.4 | 3.4 FIX antigen levels and PK parameters in HB patients grouped by mutation type

The significant correlation between FIXag and Alpha HL ($P = .004$; Table 1) prompted us to analyze the Alpha HL distribution in relation to missense variants. Individual FIXag and Alpha HL values and the overall relation ($r^2 = 0.386$; Pearson $r = 0.622$) are reported in Figure 4A.

TABLE 1 Correlation between FIX antigen and PK parameters of rFIX infused in HB patients

		Mean (95% CI)	
FIX antigen (%)		43.70 (15.91–71.49)	
		Pearson's correlation	
		r	p
Primary PK parameters			
K 1–2 (1/hrs)	0.14 (0.06–0.23)	–0.461	.012
K 2–1 (1/hrs)	0.26 (0.13–0.40)	–0.469	.010
V1 (dL/kg)	0.93 (0.78–1.08)	0.327	.084
Secondary PK parameters			
Alpha HL ^a (hrs)	5.11 (3.61–6.61)	0.516	.004
AUC (U.h/dL)	1813 (1539–2086)	–0.107	.580
Beta HL (hrs)	43.24 (32.93–53.55)	0.397	.033
CL (dL/hrs/kg)	0.031 (0.026–0.037)	0.024	.900
CLD2 (dL/hrs/kg)	0.10 (0.04–0.16)	–0.397	.033
C _{max} (IU/dL)	58.71 (48.73–68.69)	–0.407	.029
MRT (hrs)	53.50 (42.99–64.01)	0.311	.101
V2 (dL/kg)	0.59 (0.39–0.79)	0.273	.151
V _{ss} (dL/kg)	1.52 (1.22–1.82)	0.340	.072

Note: r, Pearson's coefficient, p, Pearson analysis; numbers in bold, significant values ($P < 0.05$).

K 1–2, transfer rate from central (1) to peripheral (2) compartment; K 2–1, transfer rate from peripheral (2) to central (1) compartment; V1, Volume of central compartment.

Abbreviations: Alpha HL, alpha distribution half-life; AUC, area under the curve; Beta HL, beta elimination half-life; CI, confidence interval; CL, clearance; CLD2, inter-compartment clearance; C_{max}, at zero time extrapolated factor VIII concentration; FIX, factor IX; HB, hemophilia B; MRT, mean residence time; PK, pharmacokinetic; rFIX, recombinant factor IX; V2, volume of peripheral compartment; V_{ss}, volume of distribution at steady state.

^aNormally distributed variable.

FIX activity and antigen levels (Table 2, first line) in patients ($n = 21$) with missense mutations were higher than in patients ($n = 8$) with null (nonsense, splicing, and deletion) mutations ($P = .049$). Patients with mutations at the activation sites ($n = 7$) showed FIXag levels higher than those with null mutations ($P = .002$) and than those patients ($n = 12$) with other missense mutations (“no activation sites,” $n = 14$; $P = .001$).

The distribution of PK parameters according to the 2CP model was compared in patients with different mutation types (Table 2). Whereas differences between missense and null mutations were not detectable, missense changes at the activation site were associated with significantly lower K 1–2, K 2–1, and CLD2 values, and longer Alpha HL, Beta HL, and MRT than those observed in patients with null mutations, as well as in patients with other mutations ($n = 22$, Table 2, last column).

PK parameter values for Alpha HL, Beta HL, and MRT in HB patients classified by mutation types are reported in Figure 4B. The distribution of PK values indicated high variability, either within patients with identical F9 mutations (p.R191C, p.R226Q, p.G236D; Figure 4B) or within a single group of mutations. This observation is exemplified by the Alpha HL values of the p.R379X among null mutations, and by the p.N138H and p.A37D among missense changes not located at activation sites (Figure 4B).

3.5 | Predictors of Alpha HL, Beta HL, and MRT: FIX antigen levels and F9 genotypes

The regression analysis of predictors of Alpha HL, Beta HL, and MRT was conducted including the FIXag levels and the F9 mutation types, grouped either in missense and null, or activation site variants and other mutations (Table 3). In the model including missense versus null mutations, FIXag values were independent predictors of Alpha HL (β coefficient 0.492, $P = .011$), and as a trend of Beta HL (β coefficient 0.374, $P = .064$). In the model including activation site variants versus other mutations, F9 genotypes were independent predictors of Alpha HL and MRT, and particularly of Beta HL (β coefficient 0.699, $P = .004$).

4 | DISCUSSION

Our data provide experimental evidence, based on *in vivo* and recombinant studies, to interpret the HB mutational patterns and phenotypes associated with the recurrent substitutions at the FIX activation sites.

Substitutions at R191 and R226 differentially affected residual antigen levels, with normal or increased protein levels associated

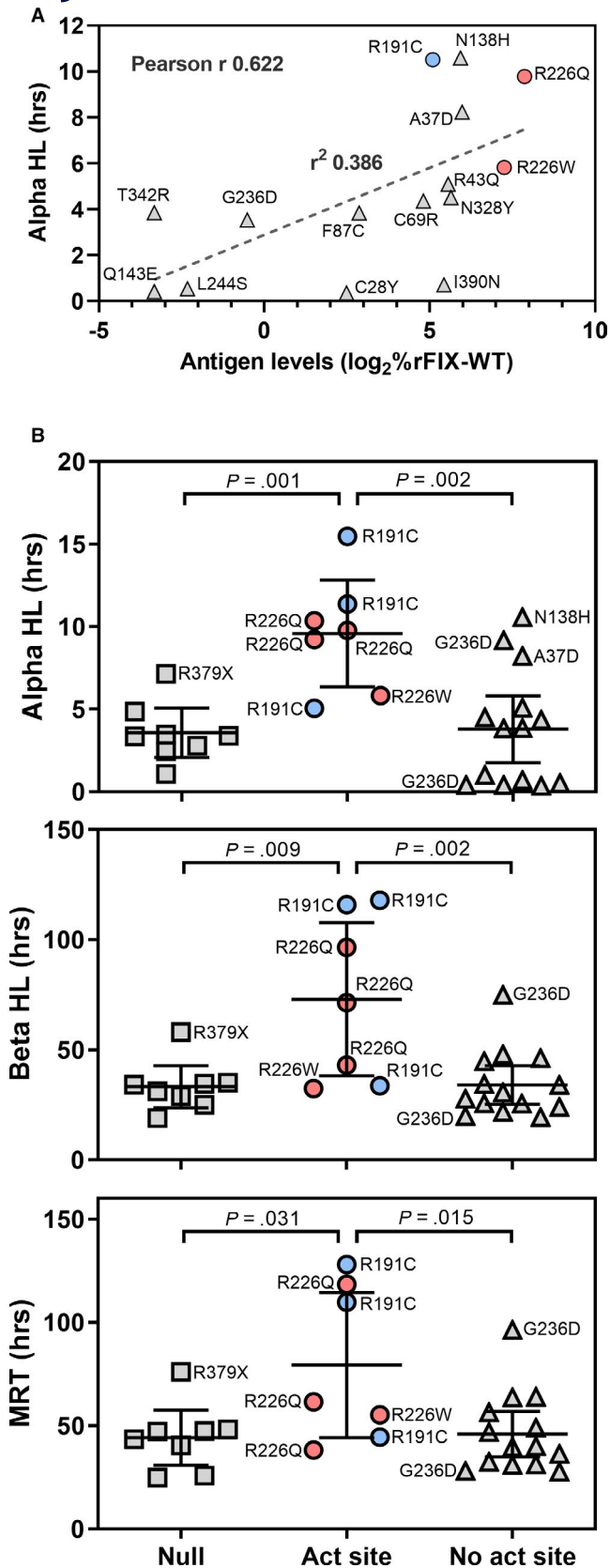


FIGURE 4 Factor IX (FIX) pharmacokinetic (PK) parameters, FIX antigen, and *F9* genotypes. A, Alpha HL parameter, *F9* genotypes and FIX antigen levels. Relation between mean antigen levels of expressed FIX missense variants of the hemophilia B (HB) cohort and Alpha HL parameter. Alpha HL of the R191C ($n = 3$ patients), R226Q ($n = 3$), and G236D ($n = 2$) variants is reported as mean of PK values from each patient. B, Alpha HL, Beta HL, and mean residence time (MRT) and *F9* genotypes. Comparative plot of selected PK parameters (Alpha HL, upper panel; Beta HL, middle panel; MRT, lower panel) among FIX variants categorized by three mutation groups: null (nonsense, splicing, deletion), missense variants affecting (act. sites) or not (no act. sites) the FIX R191 or R226 activation sites. Circles, activation site variants (blue, R191C; red, R226Q/W); gray squares, nonsense variants; gray triangles, no activation site variants

catalytically competent FIXa and cause severe HB, the benign R226K substitution, designed to mimic the positively charged WT arginine residue, permitted both efficient secretion and activation/activity, and accordingly it is not included among HB mutations. Differently, most substitutions (L/A/G) affecting the R226 position cause moderate/mild HB (Figure S2).

Multiple R191 variants are characterized by normal or modestly reduced antigen levels, and by a positive relation between activity and activity/antigen ratio. These observations suggest that the detrimental effects of substitutions at R191 on folding/secretion/scavenging processes,^{38,40,58} which all contribute to decrease antigen levels, also decrease activation and/or catalytic activity of the FIX R191 variants in a roughly proportional manner. In turn, these features shape the plasma and bleeding phenotypes of patients, and produce the graded moderate to mild HB severity of the R191C>L>P>H substitutions. The R191H change results in the mildest and most frequent form (>100 HB patients), based on its normal FIXag levels and modest activity/antigen ratio. On the other hand, the benign R191Q/W/K changes, which predict FIX activity above the levels defining mild HB,⁵⁹ are not reported in the HB nor in exome sequencing databases.

Overall, the recombinant experiments (1) assist the interpretation of the wide and graded differences in plasma FIX activity and antigen levels observed in the numerous patients affected by mutations at the R191 position, and of the degree of the associated disease severity, and (2) may predict the absence in the HB mutation database of several benign substitutions at the R191/R226 positions. A limitation of the recombinant approach is that antigen levels *in vitro* might not parallel those *in vivo*, caused by variable transfection efficiency or limited cellular capacity for post-translational modifications. However, each protein variant was expressed in six independent replicates in a cellular system widely used for its efficiency. Noticeably, the recombinant approach permits us to detect FIX variants that are secreted, and thus could bind vascular and extravascular receptors, but might have increased clearance and thus show lower FIX antigen levels in plasma (i.e., R191L). Further, the recombinant expression of FIX variants corresponding to HB causative mutations permitted us to refine the 2CP model PK analysis.

with the R226 variants. High antigen values may be also associated with other catalytically inert variants as well as with missense mutations affecting the V227 and V228 positions (Figure S2). Whereas the natural R226 substitutions do not permit the formation of a

TABLE 2 Factor IX antigen and 2CP PK parameters in HB patients grouped for F9 mutation types

Mutation types Patient number	Missense (all) 21		Null 8		Missense (activation sites) 7		Missense (no activation sites) 14	
	Mean (95% CI)	p	Mean (95% CI)	p	Mean (95% CI)	p	Mean (95% CI)	p#
FIX antigen (%)	60.05 (23.48–96.62)	.049	0.79 (–0.94–2.52)	.002	163.3 (43.97–228.5)	.001	21.95 (7.30–36.60)	<.0001
PK parameters								
K 1–2 (1/hrs)	0.17 (0.05–0.29)	.825	0.09 (0.01–0.16)	.004	0.03 (0.02–0.04)	.028	0.23 (0.06–0.41)	.016
K 2–1 (1/hrs)	0.30 (0.11–0.48)	.664	0.17 (0.09–0.26)	.001	0.04 (0.01–0.07)	.002	0.42 (0.16–0.68)	.001
V1 (dL/kg)	0.91 (0.78–1.04)	.981	0.99 (0.46–1.52)	.320	1.11 (0.85–1.38)	.018	0.81 (0.68–0.94)	.071
Alpha HL ^a (hrs)	5.70 (3.70–7.70)	.196	3.56 (2.07–5.06)	.001	9.53 (6.32–12.74)	.002	3.79 (1.76–5.81)	<.0001
AUC (U.h/dL)	1857 (1526–2189)	.633	1696 (1089–2303)	.966	1757 (988–2527)	.576	1907 (1503–2311)	.717
Beta HL (hrs)	47.04 (33.18–60.90)	.299	33.28 (23.73–42.82)	.009	72.97 (38.15–107.8)	.002	34.07 (25.33–42.82)	<.0001
CL (dL/hrs/kg)	0.030 (0.024–0.035)	.438	0.036 (0.019–0.053)	.701	0.032 (0.017–0.047)	.778	0.029 (0.023–0.035)	.978
CLD2 (dL/hrs/kg)	0.12 (0.04–0.20)	.836	0.07 (0.04–0.10)	.008	0.03 (0.02–0.04)	.068	0.16 (0.04–0.28)	.042
Cmax (IU/dL)	58.35 (45.84–70.85)	.866	59.66 (39.29–80.05)	.092	42.60 (30.41–54.73)	.023	66.22 (49.09–83.35)	.021
MRT (hrs)	57.06 (43.17–70.95)	.297	44.16 (30.84–57.49)	.031	79.39 (44.26–114.51)	.015	45.90 (34.87–56.93)	.005
V2 (dL/kg)	0.64 (0.37–0.92)	.674	0.45 (0.31–0.59)	.050	1.14 (0.35–1.92)	.006	0.39 (0.28–0.50)	.003
Vss (dL/kg)	1.55 (1.17–1.93)	.756	1.44 (0.81–2.07)	.087	2.25 (1.25–3.25)	.004	1.20 (1.00–1.40)	.004

Note: PK parameters in F9 mutation type groups were compared by *t*-test; the table includes four columns reporting *p* values, three of them flanked by the parameter values subjected to statistical evaluation. For *p*#, the comparison is made between missense mutations in the activation sites (*n* = 7) and other mutations (the sum of null, missense no activation site, *n* = 22; parameter values not reported in the table); numbers in bold, significant values (*P* < 0.05).

Abbreviations: Alpha HL, alpha distribution half-life; AUC, area under the curve; Beta HL, beta elimination half-life; CI, confidence interval; CL, clearance; CLD2, inter-compartment clearance; Cmax, at zero time extrapolated factor VIII concentration; FIX, factor IX; HB, hemophilia B; MRT, mean residence time; PK, pharmacokinetic; rFIX, recombinant factor IX; V2, volume of peripheral compartment; Vss, volume of distribution at steady state.

^aNormally distributed variable. PK parameters definition as in Table 1.

TABLE 3 Linear regression models for predictors of the PK parameter variability

	β -coefficient	<i>p</i>	Predictors
Alpha HL (hrs)	0.492	.011	FIX antigen
	-0.066	.719	<i>F9</i> genotypes—Missense vs. null mutations
	0.102	.642	FIX antigen
	0.569	.015	<i>F9</i> genotypes—Activation sites vs. other mutations
Beta HL (hrs)	0.374	.064	FIX antigen
	-0.062	.751	<i>F9</i> genotypes—Missense vs. null mutations
	-0.111	.623	FIX antigen
	0.699	.004	<i>F9</i> genotypes—Activation sites vs. other mutations
MRT (hrs)	0.274	.182	FIX antigen
	-0.099	.623	<i>F9</i> genotypes—Missense vs. null mutations
	-0.121	.625	FIX antigen
	0.594	.023	<i>F9</i> genotypes—Activation sites vs. other mutations

Note: Definition of PK parameters and mutation types as in Tables 1 and 2, respectively. The FIX antigen and PK parameter values in *F9* mutation groups are reported in Table 2. Missense mutations (patients $n = 21$) vs. null mutations (patients $n = 8$); mutations in the activation sites (patients $n = 7$) vs. other mutations (patients $n = 22$). β -coefficients, standardized regression coefficients; numbers in bold, significant values ($P < 0.05$). For each PK parameter (Alpha HL, Beta HL and MRT), the results from the linear regression models are separated by a continuous line.

Abbreviations: Alpha HL, alpha distribution half-life; Beta HL, beta elimination half-life; FIX, factor IX; MRT, mean residence time; PK, pharmacokinetic.

Previous observations in HB patients led us to hypothesize that the FIX CRM⁺ status may favorably influence^{33,46} the PK of infused FIX or that dysfunctional endogenous FIX would impair prophylaxis in HB mouse models.³⁵ With potentially discrepant information as background, we investigated in HB patients the hypothesis that dysfunctional endogenous FIX, which circulates in variable amounts, might positively or negatively modulate the distribution of infused FIX. Prompted by the presence in our HB cohort of several patients affected by mutations at the activation sites causing retention of the activation peptide, previously reported to act as a plasma retention signal,^{31,32} we also explored whether the variably abundant antigen levels in these patients could further modulate the persistence of infused FIX in plasma. It is worth noting that our study is the first report on the relation between FIX PK parameters and specific *F9* mutations and well-defined mutation groups.

The consistent correlation of primary and secondary PK parameters with dysfunctional endogenous FIX levels in HB patients supported the study of patients grouped by mutation types, which in turn dictates the concentration and features of dysfunctional molecules. Although FIXag levels were higher in patients with missense mutations than in those with null ones, their mean PK parameter values did not significantly differ, which may be explained by very low or null antigen levels associated to several missense changes (Figure 4A). The wide variability in antigen levels for the non-activation site missense changes could contribute to obscure mutation group differences in PK parameters. The grouping of missense and null mutations in linear regression analysis indicated FIXag levels, but not *F9* genotypes, as significant predictors of PK parameters, particularly of Alpha HL, whereas the activation site variants were associated with longer Alpha HL, Beta HL, and MRT than null mutations. The MRT, a parameter that may influence patients'

treatment, was 80% longer (79.39 h, 95% CI 44.3–114.5) than in patients with null mutations (44.2 h, 95% CI, 30.8–57.5, $P = .031$) or with other missense changes (45.9 h, 95% CI 34.9–56.9, $P = .015$). The regression analysis suggested activation site variants as independent predictors of several PK parameters, and particularly of Beta HL (β coefficient 0.699, $P = .004$), which may be prolonged by the increasing proportion over time of the abundant endogenous FIX variants compared to infused FIX. The very high endogenous/infused FIX ratio reached in the Beta elimination phase could magnify the effects of activation site variants, and the increasing competition for receptors may decrease both the forward and backward flow between central plasma and extravascular compartments^{38–40} of infused FIX (Figure S5 in supporting information), thus prolonging its Alpha and Beta half-lives. The contribution to FIX PK of missense mutations, potentially produced also by other mutation types,⁶⁰ may consist of quantitative components such as the residual FIXag levels, and in addition, for the R226 and R191 substitutions, of qualitative components such as the retention of the activation peptide cleaved at only one site. It has been demonstrated that removal of the activation peptide from zymogen FIX results in exposure of a binding site for low-density lipoprotein receptor-related protein, which favors FIXa catabolism.⁵⁸ Whereas mutations at position R191 may modulate PK outcomes in the presence of intermediate FIXag levels, those at position R226 may influence PK parameters in the presence of high levels of the secreted FIX variants. It is tentative to speculate that in female carriers both levels of the FIX R226W variant and FIX WT would be increased, as suggested by the increased FIXag levels and the half-reduced activity/antigen ratio.

Although based on a limited number of patients and a small number of related female carriers, data suggest that mutations at the activation sites have pleiotropic effects in plasma by

dictating in patients (1) the residual FIX activity (only variants at R191); (2) the variably abundant FIXag; and by potentially modulating (3) the PK features of the infused FIX and; in female carriers, (4) the variant and wild-type FIX levels. These *F9* genetic components, together with others producing large differences among patients affected by identical *F9* mutations, are expected to contribute to the wide and elusive variability of PK outcomes in HB patients.

Our data support, in patients characterized for *F9* mutations, the hypothesis³³ that the FIX CRM⁺ status might partially improve PK of FIX infused in HB patients, and that the activation peptide, retained in the activation site variants, may contribute to prolong the Beta phase of FIX distribution in plasma, as inferred in mouse models.^{31,32} The relative impacts of FIXag levels versus retained activation peptide on FIX PK were indirectly disentangled by linear regression analysis that suggested activation site mutations as independent predictors, which warrants further studies. These *F9*-related features would permit us to better estimate the improved pharmacokinetic profile of currently used and novel long-acting FIX molecules.^{61,62}

5 | CONCLUSIONS

In vivo and recombinant results suggest that combination of FIXag levels and type of dysfunctional molecules predicted in plasma by *F9* mutations, particularly those at the activation sites, may exert pleiotropic effects on (1) the endogenous FIX and thus HB patients' phenotypes, as well as (2) infused FIX half-lives and thus MRT in substitutive treatment.

ACKNOWLEDGMENTS

This study was financially supported by an unconditioned Investigator Initiated Research grant from Pfizer (project WI225911). The authors would like to thank Dr. Tommaso Zordan and Dr. Martina Bonsi for their help in the production of expression vectors.

CONFLICTS OF INTEREST

A.B. reports grants and personal fees from Pfizer, grants from Bayer, and grants and non-financial support from Grifols, outside the submitted work. M.M. reports personal fees from Kedrion, Novo Nordisk, SOBI, Bayer, Bioverativ, Octapharma, CSL Behring and grants from Pfizer, outside the submitted work. A.C.M. has acted as a paid consultant to Bayer, CSL, Kedrion, Novo Nordisk, Pfizer, Roche, Shire, and Sobi, and as a paid invited speaker for Bayer, CSL, Novo Nordisk, Shire, and Sobi, outside the submitted work. G.C. reports personal fees from Roche, Bayer, Shire, Uniqure, Kedrion, Novo Nordisk, and Werfen; grants and personal fees from CSL Behring and Sobi; and grants from Pfizer, outside the submitted work. M.P. received grants from Novo Nordisk and Pfizer and personal fees from Pfizer, outside the submitted work. F.B. reports grants from Bayer, outside the submitted work, and the financial support of GePKHIS by an unconditioned Investigator Initiated Research grant

from Pfizer. The remaining authors state that they have no conflicts of interest to declare.

AUTHOR CONTRIBUTIONS

A.B. created recombinant vectors and performed expression studies on FIX recombinant variants; M.M. performed PK analysis; B.L. designed and performed statistical analysis of PK analysis data; A.B. and B.L. performed statistical analysis of recombinant expression and of *F9* genotype-PK parameters' association; D.B., P.R., L.B., M.L.S., P.G., D.C., A.C.M., M.S.N., and G.C. characterized and recruited patients, collected plasma samples according to the PK study design, and performed clotting assays; E.B. provided assistance on the pharmacovigilance study; M.P. and G.C. critically revised the manuscript; A.B., M.M., and F.B. conceived the study and designed research; A.B., M.M., B.L., and F.B. analyzed and interpreted data and wrote the manuscript. All authors revised and approved the final version of the manuscript.

ORCID

Alessio Branchini  <https://orcid.org/0000-0002-6113-2694>

Massimo Morfini  <https://orcid.org/0000-0001-6565-4943>

Barbara Lunghi  <https://orcid.org/0000-0002-1517-4762>

Loredana Bury  <https://orcid.org/0000-0003-0333-0948>

Paola Giordano  <https://orcid.org/0000-0002-7762-3500>

Angelo Claudio Molinari  <https://orcid.org/0000-0002-8078-8402>

Mariasanta Napolitano  <https://orcid.org/0000-0001-8934-2936>

Elisabetta Bigagli  <https://orcid.org/0000-0003-1892-4343>

Giancarlo Castaman  <https://orcid.org/0000-0003-4973-1317>

Mirko Pinotti  <https://orcid.org/0000-0002-4114-7055>

Francesco Bernardi  <https://orcid.org/0000-0003-0305-2212>

REFERENCES

1. Monroe DM, Hoffman M. What does it take to make the perfect clot? *Arterioscler Thromb Vasc Biol.* 2006;26:41-48.
2. Lesk AM, Fordham WD. Conservation and variability in the structures of serine proteinases of the chymotrypsin family. *J Mol Biol.* 1996;258:501-537.
3. Yousef GM, Elliott MB, Kopolovic AD, Serry E, Diamandis EP. Sequence and evolutionary analysis of the human trypsin subfamily of serine peptidases. *Biochim Biophys Acta.* 2004;1698:77-86.
4. Di Scipio RG, Kurachi K, Davie EW. Activation of human factor IX (Christmas factor). *J Clin Invest.* 1978;61:1528-1538.
5. Bajaj SP, Rapaport SI, Russell WA. Redetermination of the rate-limiting step in the activation of factor IX by factor XIa and by factor VIIIa/tissue factor. Explanation for different electrophoretic radioactivity profiles obtained on activation of 3H- and 125I-labeled factor IX. *Biochemistry.* 1983;22:4047-4053.
6. Lawson JH, Mann KG. Cooperative activation of human factor IX by the human extrinsic pathway of blood coagulation. *J Biol Chem.* 1991;266:11317-11327.
7. Geng Y, Verhamme IM, Messer A, et al. A sequential mechanism for exosite-mediated factor IX activation by factor XIa. *J Biol Chem.* 2012;287:38200-38209.
8. Koeberl DD, Bottema CD, Ketterling RP, Bridge PJ, Lillicrap DP, Sommer SS. Mutations causing hemophilia B: direct estimate of the underlying rates of spontaneous germ-line transitions,

- transversions, and deletions in a human gene. *Am J Hum Genet.* 1990;47:202-217.
9. Rallapalli PM, Kemball-Cook G, Tuddenham EG, Gomez K, Perkins SJ. An interactive mutation database for human coagulation factor IX provides novel insights into the phenotypes and genetics of hemophilia B. *J Thromb Haemost.* 2013;11:1329-1340.
 10. Lenting PJ, ter Maat H, Clijsters PP, Donath MJ, van Mourik JA, Mertens K. Cleavage at arginine 145 in human blood coagulation factor IX converts the zymogen into a factor VIII binding enzyme. *J Biol Chem.* 1995;270:14884-14890.
 11. Zögg T, Brandstetter H. Activation mechanisms of coagulation factor IX. *Biol Chem.* 2009;390:391-400.
 12. Gailani D, Geng Y, Verhamme I, et al. The mechanism underlying activation of factor IX by factor XIa. *Thromb Res.* 2014;133(Suppl 1):S48-S51.
 13. Green PM, Montandon AJ, Ljung R, et al. Haemophilia B mutations in a complete Swedish population sample: a test of new strategy for the genetic counselling of diseases with high mutational heterogeneity. *Br J Haematol.* 1991;78:390-397.
 14. Usharani P, Warn-Cramer BJ, Kasper CK, Bajaj SP. Characterization of three abnormal factor IX variants (Bm Lake Elsinore, Long Beach, and Los Angeles) of hemophilia-B. Evidence for defects affecting the latent catalytic site. *J Clin Invest.* 1985;75:76-83.
 15. Diuguid DL, Rabiet MJ, Furie BC, Furie B. Molecular defects of factor IX Chicago-2 (Arg 145--His) and prothrombin Madrid (Arg 271--Cys): arginine mutations that preclude zymogen activation. *Blood.* 1989;74:193-200.
 16. Liddell MB, Peake IR, Taylor SA, Lillcrap DP, Giddings JC, Bloom AL. Factor IX Cardiff: a variant factor IX protein that shows abnormal activation is caused by an arginine to cysteine substitution at position 145. *Br J Haematol.* 1989;72:556-560.
 17. Chen SH, Schoof JM, Weinmann AF, Thompson AR. Heteroduplex screening for molecular defects in factor IX genes from hemophilia B families. *Br J Haematol.* 1995;89:409-412.
 18. Nielsen LR, Scheibel E, Ingerslev J, Schwartz M. Detection of ten new mutations by screening the gene encoding factor IX of Danish hemophilia B patients. *Thromb Haemost.* 1995;73:774-778.
 19. Attali O, Vinciguerra C, Trzeciak MC, et al. Factor IX gene analysis in 70 unrelated patients with hemophilia B: description of 13 new mutations. *Thromb Haemost.* 1999;82:1437-1442.
 20. Belvini D, Salviato R, Radossi P, et al. Molecular genotyping of the Italian cohort of patients with hemophilia B. *Haematologica.* 2005;90:635-642.
 21. Halldén C, Mårtensson A, Nilsson D, et al. Origin of Swedish hemophilia B mutations. *J Thromb Haemost.* 2013;11:2001-2008.
 22. Huang MN, Kasper CK, Roberts HR, Stafford DW, High KA. Molecular defect in factor IX Hilo, a hemophilia Bm variant: Arg--Gln at the carboxyterminal cleavage site of the activation peptide. *Blood.* 1989;73:718-721.
 23. Lin SW, Shen MC. Genetic basis and carrier detection of hemophilia B of Chinese origin. *Thromb Haemost.* 1993;69:247-252.
 24. Monroe DM, McCord DM, Huang MN, et al. Functional consequences of an arginine180 to glutamine mutation in factor IX Hilo. *Blood.* 1989;73:1540-1544.
 25. Suehiro K, Kawabata S, Miyata T, et al. Blood clotting factor IX BM Nagoya. Substitution of arginine 180 by tryptophan and its activation by alpha-chymotrypsin and rat mast cell chymase. *J Biol Chem.* 1989;264:21257-21265.
 26. Solera J, Magallón M, Martín-Villar J, Coloma A. Identification of a new hemophilia BM case produced by a mutation located at the carboxy terminal cleavage site of activation peptide. *Br J Haematol.* 1991;78:385-389.
 27. Wulff K, Bykowska K, Lopaciuk S, Herrmann FH. Molecular analysis of hemophilia B in Poland: 12 novel mutations of the factor IX gene. *Acta Biochim Pol.* 1999;46:721-726.
 28. Onay UV, Kavakli K, Kiliç Y, et al. Molecular pathology of hemophilia B in Turkish patients: identification of a large deletion and 33 independent point mutations. *Br J Haematol.* 2003;120:656-659.
 29. Hougie C, Twomey JJ. Haemophilia Bm: a new type of factor-IX deficiency. *Lancet.* 1967;1:698-700.
 30. Sabatino DE, Armstrong E, Edmonson S, et al. Novel hemophilia B mouse models exhibiting a range of mutations in the factor IX gene. *Blood.* 2004;104:2767-2774.
 31. Johansson L, Karpf DM, Hansen L, Pelzer H, Persson E. Activation peptides prolong the murine plasma half-life of human factor VII. *Blood.* 2011;117:3445-3452.
 32. Begbie ME, Mamdani A, Gataiance S, et al. An important role for the activation peptide domain in controlling factor IX levels in the blood of hemophilia B mice. *Thromb Haemost.* 2005;94:1138-1147.
 33. Roth DA, Kessler CM, Pasi KJ, et al. Human recombinant factor IX: safety and efficacy studies in hemophilia B patients previously treated with plasma-derived factor IX concentrates. *Blood.* 2001;98:3600-3606.
 34. Jin D-Y, Zhang T-P, Gui T, Stafford DW, Monahan PE. Creation of a mouse expressing defective human factor IX. *Blood.* 2004;104:1733-1739.
 35. Cooley B, Broze GJ, Mann DM, Lin F-C, Pedersen LG, Stafford DW. Dysfunctional endogenous FIX impairs prophylaxis in a mouse hemophilia B model. *Blood.* 2019;133:2445-2451.
 36. Mann DM, Stafford KA, Poon M-C, Matino D, Stafford DW. The function of extravascular coagulation factor IX in haemostasis. *Haemophilia.* 2021;27(3):332-339.
 37. Smith KJ, Thompson AR. Labeled factor IX kinetics in patients with hemophilia-B. *Blood.* 1981;58:625-629.
 38. Cheung WF, van den Born J, Kühn K, Kjellén L, Hudson BG, Stafford DW. Identification of the endothelial cell binding site for factor IX. *Proc Natl Acad Sci USA.* 1996;93:11068-11073.
 39. Cooley B, Funkhouser W, Monroe D, et al. Prophylactic efficacy of BeneFIX vs Alprolix in hemophilia B mice. *Blood.* 2016;128:286-292.
 40. Stafford DW. Extravascular FIX and coagulation. *Thromb J.* 2016;14:35.
 41. Goodeve AC, Reitsma PH, McVey JH, Working Group on Nomenclature of the Scientific and Standardisation Committee of the International Society on Thrombosis and Haemostasis. Nomenclature of genetic variants in hemostasis. *J Thromb Haemost.* 2011;9:852-855.
 42. Pignani S, Todaro A, Ferrarese M, et al. The chaperone-like sodium phenylbutyrate improves factor IX intracellular trafficking and activity impaired by the frequent p. R294Q mutation. *J Thromb Haemost.* 2018;16:2035-2043.
 43. Branchini A, Ferrarese M, Campioni M, et al. Specific factor IX mRNA and protein features favor drug-induced readthrough over recurrent nonsense mutations. *Blood.* 2017;129:2303-2307.
 44. Ferrarese M, Testa MF, Balestra D, Bernardi F, Pinotti M, Branchini A. Secretion of wild-type factor IX upon readthrough over F9 prepeptide nonsense mutations causing hemophilia B. *Hum Mutat.* 2018;39:702-708.
 45. Emsley J, McEwan PA, Gailani D. Structure and function of factor XI. *Blood.* 2010;115:2569-2577.
 46. Morfini M, Dragani A, Paladino E, et al. Correlation between FIX genotype and pharmacokinetics of Nonacog alpha according to a multicentre Italian study. *Haemophilia.* 2016;22:537-542.
 47. Iorio A, Fischer K, Blanchette V, et al. Tailoring treatment of hemophilia B: accounting for the distribution and clearance of standard and extended half-life FIX concentrates. *Thromb Haemost.* 2017;117:1023-1030.
 48. Lunghi B, Bernardi F, Martinelli N, et al. Functional polymorphisms in the LDLR and pharmacokinetics of factor VIII concentrates. *J Thromb Haemost.* 2019;17:1288-1296.
 49. Gabrielsson J, Weiner D. *Pharmacokinetic and Pharmacodynamic data analysis, Concepts and Application*, 4th edn. Swedish Pharmaceutical Press; 2007.

50. Björkman S, Blanchette VS, Fischer K, et al. Comparative pharmacokinetics of plasma- and albumin-free recombinant factor VIII in children and adults: the influence of blood sampling schedule on observed age-related differences and implications for dose tailoring. *J Thromb Haemost.* 2010;8:730-736.
51. Prowse CV. In vivo recovery of factor VIII following transfusion: a survey of recent data and publications to assess the influence of standards used for potency assignment. On behalf of the subcommittee on factor VIII and IX of the scientific and standardization committee of the ISTH. *Thromb Haemost.* 1995;74:1191-1196.
52. Björkman S, Folkesson A, Berntorp E. In vivo recovery of factor VIII and factor IX: intra- and interindividual variance in a clinical setting. *Haemophilia.* 2007;13:2-8.
53. Taylor SA, Liddell MB, Peake IR, Bloom AL, Lillicrap DP. A mutation adjacent to the beta cleavage site of factor IX (valine 182 to leucine) results in mild haemophilia Bm. *Br J Haematol.* 1990;75:217-221.
54. Sakai T, Yoshioka A, Yamamoto K, et al. Blood clotting factor IX Kashiwara: amino acid substitution of valine-182 by phenylalanine. *J Biochem.* 1989;105:756-759.
55. Maekawa H, Sugo T, Yamashita N, et al. Molecular defect in factor IX Tokyo: substitution of valine-182 by alanine at position P2' in the second cleavage site by factor XIa resulting in impaired activation. *Biochemistry.* 1993;32:6146-6151.
56. Sharathkumar A, Hardesty B, Greist A, et al. Variability in bleeding phenotype in Amish carriers of haemophilia B with the 31008 C->T mutation. *Haemophilia.* 2009;15:91-100.
57. Shinozawa K, Amano K, Hagiwara T, et al. Genetic analysis of carrier status in female members of Japanese hemophilia families. *J Thromb Haemost.* 2021;19:1493-1505.
58. Neels JG, van Den Berg BM, Mertens K, et al. Activation of factor IX zymogen results in exposure of a binding site for low-density lipoprotein receptor-related protein. *Blood.* 2000;96:3459-3465.
59. Bolton-Maggs PHB, Pasi KJ. Haemophilias A and B. *Lancet.* 2003;361:1801-1809.
60. Hildyard C, Keeling D. Effect of age on factor IX levels in symptomatic carriers of haemophilia B Leyden. *Br J Haematol.* 2015;169:448-449.
61. You CW, Hong S-B, Kim S, et al. Safety, pharmacokinetics, and pharmacodynamics of a next-generation subcutaneously administered coagulation factor IX variant, dalcinonacog alfa, in previously treated haemophilia B patients. *J Thromb Haemost.* 2021;19: 967-975.
62. Lombardi S, Aaen KH, Nilsen J, et al. Fusion of engineered albumin with factor IX Padua extends half-life and improves coagulant activity. *Br J Haematol.* 2021;194:453-462.

SUPPORTING INFORMATION

Additional supporting information may be found in the online version of the article at the publisher's website.

How to cite this article: Branchini A, Morfini M, Lunghi B, et al; The GePKHIS Study Group of AICE. F9 missense mutations impairing factor IX activation are associated with pleiotropic plasma phenotypes. *J Thromb Haemost.* 2022;20:69–81. doi:[10.1111/jth.15552](https://doi.org/10.1111/jth.15552)

APPENDIX

GEPKHIS STUDY GROUP OF AICE

Paola Agostini: Hematology and Blood Transfusion Centre, Santa Chiara Regional Hospital, Trento, Italy. Chiara Biasioli: Haemophilia Center and Transfusion Department, Hospital of Cesena, Cesena, Italy. Teresa Maria Caimi: A De Gasperis Cardiovascular Department, Niguarda Ca' Granda Hospital, Milan, Italy. Filomena Daniele: Haemostasis and Thrombosis Center, AO L'Annunziata Hospital, Cosenza, Italy. Alfredo Dragani: Haemophilia & Thrombosis Centre, Haematology Department, Regional Hospital, Pescara, Italy. Donato Gemmati: Department of Neuroscience and Rehabilitation, University of Ferrara, Ferrara, Italy. Paolo Gresele: Department of Medicine and Surgery, University of Perugia, Perugia, Italy. Silvia Linari: Department of Oncology, Center for Bleeding Disorders, Department of Oncology, Careggi University Hospital, Florence, Italy. Gina Rossetti: Hematology and Blood Transfusion Centre, Santa Chiara Regional Hospital, Trento, Italy. Cristina Santoro: Department of Haematology, "Policlinico Umberto I" University Hospital, Rome, Italy. Rita Santoro: Centre for Haemorrhagic and Thrombotic Disorders, "Pugliese Ciaccio" Hospital, Catanzaro, Italy. Gianluca Sottilotta: Haemostasis and Thrombosis Unit, Bianchi-Melacrino-Morelli' Metropolitan Hospital, Reggio Calabria, Italy. Johanna Svahn: Thrombosis and Haemostasis Unit, Gaslini Hospital, Genova, Italy.

C. Santoro acted as paid consultant/advisor/speaker for Bayer, CSL Behring, Shire/Takeda, Novo Nordisk, Amgen, Novartis, Pfizer, Roche, and Sobi. R. Santoro acted as a member of the speaker bureau or advisory board sponsored by Bayer, CSL Behring, Roche, and Shire/Takeda. The remaining members of the AICE Study group state that they have no relevant conflicts of interest for the present study.