

Table S1 Domains and items of the data collection questionnaire administered to family members enrolled in the prenatal diagnosis program for congenital factor VII deficiency.

Domain #	Questionnaire item
1. Demographic data	
1.1	Full name, date of birth, and sex of each family member
1.2	Relationship between family members (consanguinity, degree of kinship)
1.3	Referring institution and date of referral
2. Family and bleeding history	
2.1	Known family history of bleeding disorders or coagulopathy
2.2	Prior diagnosis of factor VII deficiency in any family member
2.3	Type and frequency of bleeding episodes (easy bruising, epistaxis, gingival bleeding, menorrhagia, post-surgical or post-traumatic bleeding)
2.4	Age at onset of first bleeding episode
2.5	Prior hospitalizations or interventions related to bleeding
3. Clinical and laboratory data	
3.1	Results of coagulation screening tests (PT, aPTT, fibrinogen)
3.2	FVII activity level (%) and FVII antigen level (%)
3.3	Complete blood count and other relevant laboratory parameters
3.4	Prior treatment with replacement therapy (type, dose, frequency, and response)
3.5	Presence of comorbidities or concurrent medications
4. Obstetric and reproductive history (mothers)	
4.1	Number of previous pregnancies, gestational age at delivery, and mode of delivery
4.2	History of miscarriage, termination of pregnancy, or fetal loss
4.3	Bleeding complications during pregnancy or delivery
4.4	Prior prenatal diagnosis (method, result, and outcome)
4.5	Concurrent maternal medical conditions (e.g., malignancy)
5. Genetic and molecular data	
5.1	Variant identified in the F7 gene (nucleotide and protein designation)
5.2	Zygoty (homozygous, heterozygous, compound heterozygous)
5.3	Results of STR linkage analysis for exclusion of maternal DNA contamination
5.4	Method of prenatal sampling (chorionic villus sampling) and gestational age at sampling
6. Psychosocial and ethical data	
6.1	Family decision-making regarding continuation or termination of affected pregnancies
6.2	Awareness of recurrence risk and genetic counseling received
6.3	Informed consent documentation and ethics committee approval reference

Abbreviations: PT, prothrombin time; aPTT, activated partial thromboplastin time; FVII, factor VII; STR, short tandem repeat; F7, coagulation factor VII gene.