

LETTER TO THE EDITOR

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# Three novel *F8* mutations in sporadic haemophilia A cases

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Hemophilia A (HA) is an X-linked hereditary disorder characterized by bleeding of variable severity through mild, moderate to severe owing to large range of mutations in the Factor VIII (*F8*) gene (Bowen 2002). All kind of *F8* mutations, except repeats, have been reported for HA, in total up to 2370 (Human Genome Mutation Database 2005). A preliminary study was conducted in our lab for identification of mutations in *F8* gene in Pakistani HA patients. Correlation of *F8* mutations with clinical manifestation of HA patients was the main objective of the study. Blood samples were collected from 62 HA patients from all over the Pakistan and clinical history of all HA patients was recorded (only patients frequently visiting medical centers for the replacement of Factor VIII were selected for the study). Genomic DNA was extracted from whole blood by standard organic procedure. Specific primers (Figure 1) were designed using "Primer3" ([http://biotools.umassmed.edu/bioapps/primer3\\_www.cgi](http://biotools.umassmed.edu/bioapps/primer3_www.cgi)) to amplify the coding region of *F8* gene; amplified products were sequenced by ABI 310 and ABI 3100 sequencer (Applied Biosystems, Carlsbad, CA, USA). The sequencing results were visualized using "Chromas 2.33" software (Applied Biosystems) and mutations were detected using "BLAST" software available on the NCBI website (<http://blast.ncbi.nlm.nih.gov/Blast.cgi>). Three novel mutations (1 deletion; 2 point mutations) were detected in four sporadic HA patients, all from different ethnic backgrounds (Table 1). The deletion of T in exon 7 within the A1 domain represents a frame-shift change disrupting the protein structure and function, which result in severe manifestation of the disease. A missense point mutation in the A3 domain occurs in codon 1907 at nucleotide number 5720, replacing Serine with Isoleucine, and

confers a moderate type of severity. It should be noted that Serine is a polar and acidic amino acid while Isoleucine is a nonpolar and basic amino acid. A nonsense point-mutation was found in two unrelated patients in the C3 domain (exon 26) and was correlated with moderate clinical findings. Beside these mutations, 27 common SNPs were also detected in *F8* gene for the studied patients (Table 2). The allelic data and accession numbers of these SNPs were collected from Ensembl Genome Browser (Ensembl 2000). The results of the study will form the basis not only for an enlarged study but also for diagnosis and genetic counseling of classical hemophilia in Pakistan.

## Competing interests

The authors declare that they have no competing interests.

## Author's contributions

RH managed the project and wrote the paper. NBA, SH, ZS, MA, SA performed experiments. GN designed the project. All authors read and approved the final manuscript.

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## References

- Bowen DJ (2002) Hemophilia A and B: molecular insights. *J clin path; mol path* 55:1–18.
- Human Genome Mutation Database (2005) Institute of Medical Genetics, Cardiff, <http://www.hgmd.cf.ac.uk>. Accessed 26 May 2012.
- Ensembl (2000) European Molecular Biology Laboratory and Wellcome Trust Sanger Institute, <http://www.ensembl.org>. Accessed 10 June.

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Primer ID	Exon	Primer	Size	Product size	Annealing Temp.	FRE14.1F	14	GGGAGAGAACTCTACAGAA	21	396bp	55.18°C	FRE20R		ATCTGAGATTCTCCACAGA	20	54.6°C
FRE1F	1	CTGCTTCCACTGATAAAA	20	457bp	55.55°C	FRE14.1R		GCATCAACAATCTACTAGAG	21		54.97°C	FRE21F		CTAGGACTAACCAGCTGAA	20	54.7°C
FRE1R		AGCATCACAACCTCTTAAC	20		55.06°C	FRE14.2F	14	CCACAATTCAGAAAATGAC	20	463bp	55.47°C	FRE21R	21	GAGCTTCAAGAGGAATAAG	210bp	54.5°C
FRE2F	2	AAGTGTCACCAAAATGAAC	20	258bp	54.94°C	FRE14.3F	14	TAAATGAGAACTGGGGACA	20		55.65°C	FRE22F		TTCAGGAGTAGCACATACA	20	54.2°C
FRE2R		GGACHTTTAACTGGAACCT	20		54.80°C	FRE14.4F	14	ATGGACCTGCTTTGTGAC	19	487bp	56.01°C	FRE22R	22	AGTATTCCAGCATTCCTTT	20	55.4°C
FRE3F	3	TGGAATAACAGGTTTCTGG	20	250bp	55.23°C	FRE14.4R		TGACCTTCCACAGATTTTC	20		55.17°C	FRE23F		GCACAAAGCAATAGAAAG	20	55.3°C
FRE3R		GCACACACATCTCAGCTTC	20		54.86°C	FRE14.5F	14	GGATACAAAGGACTCATGA	20	487bp	54.95°C	FRE23R	23	GTTGAGGGAAGAGGATATG	20	53.8°C
FRE4F	4	TGTTCTTTGAGTATACATGG	22	371bp	55.49°C	FRE14.5R	14	TTTGAGAAATGAGCTGTGTG	20		54.89°C	FRE24F		GCATGCTCTGTGATACTG	20	55.1°C
FRE4R		TCTTCAGGTGAAGGAACAC	20		55.24°C	FRE14.6F	14	GGCATATGCTCCAGTACTTC	20	529bp	54.90°C	FRE24R	24	ACCTCAGAAGAACAGTCAAG	21	53.7°C
FRE5F	5	TCTCTAGTGACAAATTC	20	188bp	54.23°C	FRE14.6R	14	TGCTGGAAATGAGAAGAT	20		55.05°C	FRE25F		TCTGGAGTAAATGGTGAC	20	55.0°C
FRE5R		GCAGAGGATTTCTTTCAGG	19		54.97°C	FRE14.7F		GAGTCATAGCATCCCTCAAG	20	492bp	54.85°C	FRE25R	25	TTAAGCTCTAGGAGGTTGTT	21	55.0°C
FRE6F	6	TGCTCATGAGACATGCTG	20	231bp	54.9°C	FRE14.7R	14	CTGTTGCTTCATCCACTT	20		55.45°C	FRE26.1F		AGAAGTGAGAAAAGCGTCTG	20	54.89°C
FRE6R		ACAGAACTCTGGTCTGAAT	20		54.8°C	FRE14.8F	14	AGAAAGCACTATCCCTACG	20	366bp	55.02°C	FRE26.1R	26	GGAAAGGAGGAGTAATCTGG	20	55.38°C
FRE7F	7	TCCATCTGCTCTAGCAAGT	20	400bp	54.9°C	FRE14.8R	14	TGACCTTCTTTGGGCTTA	20		55.23°C	FRE26.2F		ATCATGCTCTGCTATTCT	20	54.66°C
FRE7R		CCTCAGCAACACTATATTC	22		54.2°C	FRE14.9F	14	GATACATTTTGGCCGTAA	20	445bp	54.97°C	FRE26.2R	26	GTCGCCCTCATAGCTAA	20	55.17°C
FRE8F	8	GCCTAATAGCAAGACATCTG	23	358bp	55.2°C	FRE14.9R	14	TGGTGTCATCATCTGGTAA	20		54.75°C	FRE26.3F		ACAATCGAAAATGGAGAG	20	55.37°C
FRE8R		TTTGAAGTATGGGAAGAGA	20		54.8°C	FRE15F	15	GAGGATGTGAGGCATTCTA	20	300bp	55.3°C	FRE26.3R	26	GGGAGAGAGTAACTGAGTGC	21	55.70°C
FRE9F	9	ATTTTCTTCCCACTCTC	20	302bp	54.9°C	FRE15R		GTTGGAAATACATATAGCTAGC	22		53.1°C	FRE26.4F		GATGACATTAGGCTTCTAAAG	22	54.95°C
FRE9R		GACAAAGGTGAATATGAGG	20		54.9°C	FRE16F	16	GGATGTAAACCTAAGGAC	20	389bp	55.1°C	FRE26.4R	26	TTAGGATCTCTGTTTCCA	20	54.85°C
FRE10F	10	GGCCACTTTATTTATCTGG	20	284bp	54.1°C	FRE16R	16	AGCTCTTATGACACTAGG	20		54.9°C	FRE26.5F		GGCTGGAGACAAGGATAAGT	20	55.90°C
FRE10R		CTGGAAAGGACCAACATA	20		55.2°C	FRE17F	17	TGAGAAATCCACTCTGGTTC	20	371bp	55.6°C	FRE26.5R	26	CAGTGCCCTATTGTTTGA	20	55.43°C
FRE11F	11	CAGATTGTAGAACCTTGC	20	361bp	55.0°C	FRE17R	17	CTGGATCAATCTCATTTG	20		55.6°C					
FRE11R		AAGGGACATACACTGAGAA	20		54.6°C	FRE18F	18	ATACTGTGGGAGTGAATC	20	389bp	53.81°C					
FRE12F	12	GACTCTAGCTCTACTGGA	20	262bp	54.8°C	FRE18R	18	TCGCTTGATCAGTATTG	20		54.79°C					
FRE12R		TCTTTATTCACCACTCTG	20		56.0°C	FRE19F	19	ACCAATGTATCTCATGCTCA	20	226bp	53.9°C					
FRE13F	13	TCTTCTCTGGGAATAAGAT	20	393bp	53.0°C	FRE19R	19	AGGCTGAGTGGTGAAGAAC	20		55.1°C					
FRE13R		ATAGGAATGCTAGTGAAGC	20		54.7°C	FRE20F	20	GCTGAATTTGTGCACTCT	20	199bp	55.6°C					

**Figure 1** Primers used in the study.

**Table 1** Novel mutations in *F8* gene

Age/Sex	Severity	Exon	Nucleotide change	Amino acid change	Codon/Codon no.	Nucleotide genome ref./cDNA ref.	Affected Domain
4 yr /male	Severe	7	Deletion of T	Frame-shift	CTC → C-C/ 318	159197688/953	A1
35 yr / male	Moderate	17	G → T	Ser → Ile	AGC → ATC/ 1907	154132724/5720	A3
15 & 19 yr /male	Moderate	26	C → A	Tyr → Termination	TAC → TAA/ 2324	154065994/6972	C2

yr (years).

**Table 2** Common SNPs in *F8* gene (exonic region)

Sr. #	Patient	Exon	SNP ambiguity	SNP	Codon	Codon#	Comments	Accession number
1	All 62 Samples	2	W: A/T	A/A	<u>G</u> AT	75	European = T/T	rs1800288
2	All 62 Samples	7	K: G/T	G/G	T <u>G</u> G	274	European = C/C; Spanish Caucasians = C(0.995)/A(0.005); African American, Chinese, Southeast Asia, Mexican Indian = C/A	rs34371500
3	All 62 Samples	8	R: G/A	G/G	<u>C</u> GC	391	Ancestral: <b>G</b>	rs137852364
4	All 62 Samples	8	Y: T/C	T/T	<u>T</u> CA	392	European = C/C	rs28933669
5	All 62 Samples	8	Y: C/T	C/C	<u>T</u> CA	392	?	rs28933668
6	All 62 Samples	8	K: T/G	T/T	<u>A</u> TT	405	European = A/A	rs28933670
7	All 62 Samples	8	R: A/G	A/A	GAG	409	?	rs28933671
8	All 62 Samples	9	K: G/T	T/T	<u>T</u> TG	431	Ancestral: <b>G</b>	rs28933672
9	All 62 Samples	9	R: A/G	A/A	<u>A</u> AA	444	Ancestral: <b>G</b>	rs28937272
10	All 62 Samples	9	W: T/A	T/T	<u>I</u> AC	450	Ancestral: <b>A</b>	rs111033616
11	All 62 Samples	10	R: G/A	G/G	<u>C</u> GT	503	Ancestral: <b>A</b>	rs35383156

**Table 2 Common SNPs in F8 gene (exonic region) (Continued)**

12	All 62 Samples	12	Y: T/C	T/T	<u>CTT</u>	622	Ancestral: <b>T</b>	rs1800290
13	All 62 samples	15	R: G/A	G/G	<u>CAG</u>	1764	Ancestral: <b>A</b>	rs5986891
14	All 62 samples	16	R: G/A	G/G	<u>ATG</u>	1842	European = G/G	rs28943674
15	All 62 samples	16	Y: C/T	C/C	<u>CCC</u>	1844	European = C/C	rs28933675
16	All 62 samples	16	M: A/C	A/A	<u>ACT</u>	1845	?	rs28933676
17	All 62 samples	16	Y: C/T	C/C	<u>GCC</u>	1853	European = C/C	rs28933677
18	All 62 samples	17	D: G/A/T	G/G	GAT	1865	Not Available	CI076951
19	All 62 samples	17	R: A/G	A/A	<u>CAC</u>	1867	Ancestral: <b>G</b>	rs28933679
20	All 62 samples	17	S: C/G	C/C	<u>CCC</u>	1873	European = G/G	rs28933680
21	All 62 samples	17	R: G/A	G/G	<u>GAG</u>	1904	European = C/C	rs28933681
22	All 62 samples	17	S: G/C	G/G	<u>TGC</u>	1922	European = G/G	rs4384155
23	All 62 samples	17	S: C/G	C/C	<u>TGC</u>	1922	European = C/C	rs4520342
24	All 62 samples	18	R: A/G	A/A	AAT	1940	?	CM083806
25	All 62 samples	18	D: G/A/T	G/G	<u>CGA</u>	1960	?	rs28937294
26	All 62 samples	18	R: G/A	G/G	GGC	1967	?	rs111033615
27	All 62 samples	24	Y: C/T	C/C	TAC	2214	Ancestral: <b>C</b>	rs1800296