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Abstract

Objective: Familial Mediterranean fever (FMF) is the most common autoinflammatory disease. Most of the identified disease-causing mutations are located on exon 10. As the number of studies about the effect of the exonal location of the mutation and its phenotypic expression is limited, we aimed to investigate whether the exonic location of the Mediterranean fever (MEFV) mutation has an effect on the clinical manifestation in patients with FMF.

Methods: Study population was derived from the main FMF registry that included 2246 patients from 15 different rheumatology clinics. We categorized the mutations according to their exon locations and retrieved the clinical and demographic information from the database.

Results: Patients having the *MEFV* mutations on exon 2 or 10 (n:1526) were divided into three subgroups according to the location of the *MEFV* mutations: Group 1 (exon 2 mutations), Group 2 (exon 10 mutations), and Group 3 (both exon 2 and exon 10 mutations). Group 2 patients were of a significantly younger age at onset, and erysipel-like erythema, arthritis, amyloidosis, and a family history of FMF were more common in this group.

Conclusion: Patients with FMF and exon 10 mutations show more severe clinical symptoms and outcome. Exon 2 mutations tend to have a better outcome.

Keywords: E148Q, exon 2, exon 10, familial Mediterranean fever, M694V

Introduction

Familial Mediterranean fever (FMF) is the most common form of autoinflammatory disease and inherited autosomal recessively. The responsible gene for FMF is located in the short arm of chromosome 16 and named *MEFV* (Mediterranean fever) (1).

So far, over 300 sequence variations have been identified in the *MEFV* gene (2). Most of the identified mutations, including M694V, M680I, M694I, and V726A, are located on exon 10, whereas some mutations such as E148Q and R202Q are located on exon 2 (1). It has been reported that there is a genotype-phenotype relation in FMF, such as a more severe disease with M694V and a milder disease course in patients carrying E148Q (3,4). Mutations located on exon 2, mostly E148Q, are commonly considered to lead to a better outcome, less frequent attacks, and amyloidosis (3, 4). Currently there are a limited number of reports addressing exonal location and its phenotypic expression in FMF. Additionally, studies on this subject provided conflicting results. Therefore, in the current study, we aimed to investigate whether the exonic location of the *MEFV* mutation has an effect on the clinical manifestation in patients with FMF.

Methods

Patients

Study population was derived from the main FMF registry that included 2246 (34.5±11.9 years, 46.7% male) patients from 15 adult rheumatology clinics located in different geographical parts of Turkey (5). The patients in this registry were diagnosed according to the Tel-Hashomer or Sheba Medical Center criteria (6, 7).

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We categorized the mutations according to their exon locations (exon 2 and 10) and retrieved the clinical and demographic information from the database. A more detailed description of the national FMF registry has been previously published (5).

The study complies with the Declaration of Helsinki and was approved by the local ethical committee of Eskişehir Osmangazi University (decision date: 27.08.2012; decision number: 2012/184).

Genetic analysis

There were 1719 patients with available genotype information, which was done by the PCR restriction fragment length polymorphism (PCR-RFLP) or the reverse hybridization assay (FMF StripAssay), depending on the laboratory of the participating clinics.

Statistical analysis

The normality was tested using the Shapiro-Wilk method. Based on the results, most of the variables showed non-normal distribution. Comparison of the continuous variables was done by the Kruskal-Wallis test, and the nominal or ordinal data were analyzed by a chi-squared test. Continuous

variables were presented as median and 25-75th percentiles, and the categorical variables are expressed in frequencies and percentages. As there were multiple groups, Bonferroni's correction (post hoc test) was used to reduce the probability of a Type I error. The adjusted critical significance level was <0.016.

Statistical comparisons were performed using the Statistical Package for Social Sciences (SPSS) version 21.0 (IBM Corp.; Armonk, NY, USA).

Results

There were 1719 patients included in the *MEFV* mutation analysis. One hundred and fifty-four out of 1719 patients did not have any detectable mutation, 39 had rare mutations located in other exons. Remaining 1526 patients (median age 31 [24 to 41] years, 48.4% male) were divided into three subgroups according to the location of the *MEFV* mutations: Group 1 (exon 2 mutations: E148Q and R202Q), Group 2 (exon 10 mutations: M694V, V726A, M680I, M694I, R761H, K695R), and Group 3 (compound heterozygous; having both exon 2 and exon 10 mutations). The distribution of the *MEFV* mutations by groups is listed in Table 1.

The comparison of three groups showed that patients with exon 2 mutations had female predominance (65.9%) when compared to Group 2 (51.7%) and Group 3 (42.9%) ($p < 0.001$). Patients with isolated exon 10 mutations (14[8-21]) were of a significantly younger age at onset when compared to Group 1 (20[13-26]) and Group 3 (19[13-25]) ($p < 0.001$). The demographic and clinical features of the patients are summarized in Table 2.

The frequency of fever, peritonitis, pleuritis, myalgia, vasculitis, chronic renal failure (CRF), and a family history of CRF was similar in all groups. But erysipelas like erythema (ELE), arthritis, amyloidosis, and a family history of FMF were more common in patients with only exon 10 mutations ($p < 0.05$) (Table 2).

Discussion

The *MEFV* gene is located on the short arm of chromosome 16 and has 10 exons. Most of the disease-causing mutations are located in the 10th exon. On the other hand, E148Q, one of the commonest mutations observed in FMF patients is located in the exon 2, together with R202Q. The mutations in exon 10 of the *MEFV* gene, such as M694V, V726A, M680I, M694I, R761H, K695R, are considered to cause FMF symptoms (1). In addition, certain mutations such as M694V are responsible for an early onset and severe outcome, including amyloidosis (5). In general, exon 2 mutations, particularly E148Q have been reported to be associated with a late onset disease and favorable outcome (1). Amyloidosis is the most serious complication of FMF and is generally linked with the M694V genotype. In a previous report, it has been shown that patients homozygous for the M694V genotype had six-fold risk of amyloidosis compared to patients with FMF carrying other *MEFV* gene mutations (5). Aside from amyloidosis, there has been interest to identify clinical features associated with severe disease. Some studies suggested that

Table 1. The detailed list of mutations by groups

<i>MEFV</i> mutations	Group 1 (n=82), (n, %)
E148Q homozygous	1 (1.2)
E148Q heterozygous	71 (86.6)
E148Q/R202Q	1 (1.2)
R202Q heterozygous	9 (11)
	Group 2 (n=1304), (n, %)
M694V homozygous	414 (31.7)
M694V heterozygous	293 (22.5)
M694V compound heterozygous	293 (22.5)
Non-M694V homozygous	80 (6.1)
Non-M694V heterozygous	127 (9.8)
Non-M694V compound heterozygous	97 (7.4)
	Group 3 (n=140), (n, %)
E148Q/M694V	93 (66.4)
E148Q/Non-M694V	35 (25)
R202Q/M694V	12 (8.6)

Table 2. Summary of the demographic and clinical features of the patients

	Group 1 (Patients with exon 2 mutations) n:82	Group 2 (Patients with exon 10 mutations) n:1304	Group 3 (Patients with exon 2 and 10 mutations) n:140	p	Bonferroni's correction (post hoc test)
Sex [M n (%)]	28 (34.1)	630 (48.3)	80 (57.1)	0.001	1-2: p=0.016 1-3: p=0.012 2-3: p=0.05
Age at baseline clinical visit [Median (%25-%75)]	26.5 (19.75-37)	24 (18-34)	27.5 (21.25-37.75)	0.114	
Age at onset of symptoms [Median (%25-%75)]	20 (13-26)	14 (8-21)	19 (13-25)	<0.001	2-3: <0.001 2-1: <0.001 1-3: 0.931
Fever	71 (86.6)	1196 (91.7)	128 (91.4)	0.274	
Peritonitis	78 (95.1)	1232 (94.5)	131 (93.6)	0.871	
Pleuritis	37 (45.1)	653 (50.1)	68 (48.6)	0.660	
ELE*	3 (3.7)	351 (26.9)	24 (17.1)	<0.001	1-2: <0.001 1-3: 0.003 2-3: 0.01
Arthritis	22 (26.8)	594 (45.6)	45 (32.1)	<0.0001	1-2: 0.01 1-3: 0.45 2-3: 0.002
Myalgia	12 (14.6)	199 (15.3)	19 (13.6)	0.863	
Vasculitis	2 (2.4)	100 (7.7)	5 (3.6)	0.490	
Amyloidosis	3 (3.7)	133 (10.2)	6 (4.3)	0.014	1-2: 0.02 1-3: 1 2-3: 0.05
CRF**	1 (1.2)	69 (5.3)	3 (2.1)	0.75	
Family history of FMF	32 (39)	780 (59.8)	73 (52.1)	<0.001	1-2: <0.001 1-3: 0.07 2-3: 0.08
Family history of amyloidosis	5 (6.1)	267 (20.5)	25 (17.9)	0.005	1-2: 0.01 1-3: 0.01 2-3: 0.51
Family history of CRF	3 (3.7)	79 (6.1)	7 (5)	0.606	

Data are given as n (%); *ELE: erysipel-like erythema; **CRF: chronic renal failure

patients with ELE had a more severe clinical course and were also related to M694V homozygosity and amyloidosis (8). In a previous study, researchers showed that amyloidosis, arthritis, family history of FMF, and presence of the M694V allele had been clustered in a severe outcome group (9). In this study, we showed that patients with isolated exon 2 mutations had a higher percentages of females, late onset of disease, lower rates of self, and a family history of amyloidosis, ELE, and arthritis compared to patients with exon 10 mutations. Taken together, our results are in line with the reports suggesting a favorable clinical course in patients with exon 2 mutations.

In our study, 86.5% of the patients in Group 1 had E148Q mutations, and the most common mutation in Group 2 was the homozygous M694V mutation (31.6%). Considered together with a milder disease course, our findings may support the notion that E148Q may be a polymorphism rather than a disease-causing mutation (10). However, there is still no consensus about whether the E148Q mutation is pathogenic or a simple polymorphism. In a study from Japan by Migita et al. (11), the E148Q mutation was found to be higher in FMF patients compared to the healthy population, and these patients had milder disease (11). There is also a possibility that mutations in the other allele may affect the pathogenicity of E148Q. On

the other hand, M694V is nearly always blamed for causing a more severe disease (5, 12). As we noted earlier, the other mutation in exon 2 is R202Q. This mutation is reported as one of the most common mutations in some geographic areas of Turkey (13-15). R202Q is also considered to be a polymorphism and is found to be positive in healthy population (16). Some data suggest that R202Q may be pathogenic when it is in homozygous pattern (16).

Current study has some limitations. One of them is the lack of data to calculate the severity score for FMF. The other one is the lack of full sequencing and therefore the absence of rare mutations in exon 2 and 10.

In conclusion, despite the limitations, we replicated that FMF patients with exon 10 mutations show more severe clinical symptoms and outcome. A close and careful monitoring of FMF patients with exon 10 mutations and their first degree relatives may help to diagnose FMF and its complications earlier. Further follow-up of patients with E148Q and R202Q mutations is needed to determine whether these are disease-causing mutations or polymorphisms, and a more detailed genetic testing with clinical confirmation is needed to make the final decision.

Ethics Committee Approval: Ethics committee approval was received for this study from the Ethics Committee of Eskişehir Osmangazi University (Decision Date: August 27, 2012; Approval No: 2012/184).

Informed Consent: Written informed consent was obtained from the patients who participated in this study.

Peer-review: Externally peer-reviewed.

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References

1. Sari I, Birlık M, Kasifoglu T. Familial Mediterranean fever: an updated review. *Eur J Rheumatol* 2014; 1: 21-3. [\[CrossRef\]](#)
2. Available from: URL: <http://fmf.igh.cnrs.fr/IS-SAID/infervers/index.php>
3. Ece A, Cakmak E, Uluca U, Kelekci S, Yolbas I, Gunes A, et al. The MEFV mutations and their clinical correlations in children with familial Mediterranean fever in southeast Turkey. *Rheumatol Int* 2014; 34: 207-12. [\[CrossRef\]](#)
4. Jarjour RA. Familial Mediterranean fever in Syrian patients: MEFV gene mutations and genotype-phenotype correlation. *Mol Biol Rep* 2010; 37: 1-5. [\[CrossRef\]](#)
5. Kasifoglu T, Bilge SY, Sari I, Solmaz D, Senel S, Emmungil H, et al. Amyloidosis and its related factors in Turkish patients with familial Mediterranean fever: a multi centre study. *Rheumatology (Oxford)* 2014; 53: 741-5. [\[CrossRef\]](#)
6. Livneh A, Langevitz P, Zemer D, Zaks N, Kees S, Lidar T, et al. Criteria for the diagnosis of familial Mediterranean fever. *Arthritis Rheum* 1997; 40: 1879-85. [\[CrossRef\]](#)
7. Pras M. Familial Mediterranean fever: from the clinical syndrome to the cloning of the pyrin gene. *Scand J Rheumatol* 1998; 27: 92-7. [\[CrossRef\]](#)
8. Lidar M, Doron A, Barzilai A, Feld O, Zaks N, Livneh A, et al. Erysipelas-like erythema as the presenting feature of familial Mediterranean fever. *J Eur Acad Dermatol Venereol* 2013; 27: 912-5. [\[CrossRef\]](#)
9. Akar S, Solmaz D, Kaşifoglu T, Yaşar Bilge S, Sari I, Gumus ZZ, et al. Different disease subtypes with distinct clinical expression in familial Mediterranean fever: results of a cluster analysis. *Rheumatology* 2016; 55: 343-6. [\[CrossRef\]](#)
10. Tchernitchko D, Legendre M, Cazeneuve C, Delahaye A, Niel F, Amselem S. The E148Q MEFV allele is not implicated in the development of familial Mediterranean fever. *Hum Mutat* 2013; 22: 339-40. [\[CrossRef\]](#)
11. Migita K, Izumi Y, Jiuchi Y, Iwanaga N, Kawahara C, Agematsu K, et al. Familial Mediterranean fever is no longer a rare disease in Japan. *Arthritis Res Ther* 2016; 30: 18: 175.
12. Cekin N, Akyurek ME, Pinarbasi E, Ozen F. MEFV mutations and their relation to major clinical symptoms of Familial Mediterranean Fever. *Gene* 2017; 30: 626: 9-13. [\[CrossRef\]](#)
13. Kilinc M, Ganiyusufoglu E, Sager H, Celik A, Olgar S, Cetin GY, et al. The report of sequence analysis on familial Mediterranean fever gene (MEFV) in South-eastern Mediterranean region (Kahramanmaraş) of Turkey. *Rheumatol Int* 2016; 36: 25-31. [\[CrossRef\]](#)
14. Sayın Kocakap DB, Günel-Özcan A, Çabuk F, En-sari C. The frequency of Familial Mediterranean fever gene mutations and genotypes at Kirikkale and comparison with the mean of regional MEFV mutation frequency of Turkey. *Mol Biol Rep* 2014; 41: 1419-26. [\[CrossRef\]](#)
15. Gunesacar R, Celik MM, Arica V, Elmacioglu S, Ozturk OH. Frequency of MEFV gene mutations in Hatay province, Mediterranean region of Turkey and report of a novel missense mutation (I247V). *Gene* 2014; 10: 546: 195-9. [\[CrossRef\]](#)
16. Yigit S, Karakus N, Tasliyurt T, Kaya SU, Bozkurt N, Kisacik B. Significance of MEFV gene R202Q polymorphism in Turkish familial Mediterranean fever patients. *Gene* 2012; 10506: 43-5. [\[CrossRef\]](#)