Detection of Dental Fluorosis-Associated Quantitative Trait Loci on Mouse Chromosomes 2 and 11

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Abstract

Systemic exposure to greater than optimal fluoride (F) can lead to dental fluorosis (DF). Parental A/J (DF-susceptible) and 129P3/J (DF-resistant) inbred mice were used for histological studies and to generate F2 progeny. Mice were treated with 0 or 50 ppm F in their drinking water for 60 days. A clinical criterion (modified Thylstrup and Fejerskov categorical scale) was used to assess the severity of DF for each individual F2 animal. Parental strains were subjected to histological examination of maturing enamel. F treatment resulted in accumulation of amelogenins in the maturing enamel of A/J mice. Quantitative trait loci (QTL) detection was performed using phenotypic extreme F2 animals genotyped for 354 single nucleotide polymorphism-based markers distributed throughout the mouse genome followed by \( \chi^2 \) analysis. Significant evidence of association was observed on chromosomes 2 and 11 for a series of consecutive markers (p < 0.0001). Further analyses were performed to examine whether the phenotypic effects were found in both male and female F2 mice or whether there was evidence for gender-specific effects. Analyses performed using the markers on chromosomes 2 and 11 which were significant in the mixed-gender mice were also significant when analyses were limited to only the male or female mice. The QTL detected on chromosomes 2 and 11 which influence the variation in response to fluorosis have their effect in mice of both genders. Finally, the QTL in both chromosomes appear to have an additive effect.

Key Words

Genetics · Dental fluorosis · Fluoride · Mouse models

Abbreviations used in this paper

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
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<tr>
<td>DF</td>
<td>dental fluorosis</td>
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<tr>
<td>EDTA</td>
<td>ethylenediaminetetraacetic acid</td>
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<td>F</td>
<td>fluoride</td>
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<td>NBF</td>
<td>neutral buffered formalin</td>
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<td>QTL</td>
<td>quantitative trait loci</td>
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<td>SNP</td>
<td>single nucleotide polymorphism</td>
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Introduction

Dental (enamel) fluorosis (DF) is an undesirable developmental defect of tooth enamel attributed to greater than optimal systemic fluoride (F) exposure during critical periods of amelogenesis. DF is characterized by increased porosity (subsurface hypomineralization) with a loss of enamel translucency and increased opacity [Fejerskov et al., 1990]. It is generally accepted that increasing DF severity correlates with increasing F exposure. However, individual variation in DF severity can exist when F exposure is relatively constant in a community [Mabelya et al., 1994; Yoder et al., 1998]. Our studies using inbred strains of mice indicate that genetic background plays a role in DF susceptibility and in F actions on bone biology [Everett et al., 2002; Vieira et al., 2005; Mousny et al., 2006; Yan et al., 2007]. The mechanism(s) that underlie the development of DF remain elusive, as well as the precise stage of amelogenesis most affected by F. Despite the ability of F to interact at a physicochemical manner with tooth enamel, the cellular target is the ameloblast [Pergolizzi et al., 1995]. In chronic F exposure, the maturation phase of enamel formation appears to be the target stage [Den Besten, 1986; Richards et al., 1986; Den Besten and Thariani, 1992]. Retention of enamel matrix proteins due to reduced removal during enamel maturation, perturbation of extracellular transport or initiation of endoplasmic reticulum stress response pathway indicate that some developmental process is possibly adversely affected by excessive F [Den Besten, 1986; Den Besten and Thariani, 1992; Matsuo et al., 1996; Den Besten et al., 2002; Kubota et al., 2005].

Material and Methods

Animals

Male and female A/J (DF-susceptible) and 129P3/J (DF-resistant) inbred mice were obtained from the Jackson Laboratory (Bar Harbor, Me., USA) at 5–6 weeks of age and were acclimated for 1 week prior to mating in reciprocal crosses to produce F1 hybrids. A panel of F2 mice was produced via sister:brother mating of F1 mice. At weaning, each F2 animal was placed on NaF provided in the drinking water at a concentration of 50 ppm F ion for 60 days. Parental A/J and 129P3/J mice at 3 weeks of age were treated with 0 or 50 ppm F as controls for examiner calibration and for histological studies. The treatment duration permitted full development of DF in the erupting incisors (fig. 1) [Everett et al., 2002]. All animals were housed in the Division of Lab Animal Medicine facility within the Dental Research Center, a fully AAALAC-accredited unit, and were maintained on a 12:12-hour light/dark cycle with an ambient temperature of 21°C. Mice were fed a constant nutrition (LabDiet® 5001; PMI Nutrition International, Richmond, Ind., USA), which contained 0.95% calcium, 0.67% phosphorus, 4.5 IU/g vitamin D3 and an average F of 6.56 ± 0.28 µg/g. All experimental procedures were approved by the Institutional Animal Care and Use Committee at the University of North Carolina at Chapel Hill.

Sample Collection

Serum was collected from each mouse following euthanasia and then frozen at −80°C. Samples were randomly selected for F determinations. Microdirect analysis of serum F was performed according to the method of Vogel et al. [1990].

DF Phenotyping

All mice were uniquely identified, and the 2 examiners (E.T.E. and D.Y.) performed independent assessment of DF status for each animal. The determination of DF was made clinically over the entire lower incisor tooth surfaces according to a modified Thylstrup and Fejerskov (TF) index (table 1) [Thylstrup and Fejerskov, 1978; Fejerskov et al., 1994; Everett et al., 2002].
Elite ABC kit (Rabbit IgG; Vector Laboratories, Burlingame, Calif., USA) according to the manufacturer’s instructions. Sections were counterstained using hematoxylin prior to examination.

Statistical Analyses

$\chi^2$ analysis was performed to compare the genotypic distribution in the 2 groups of phenotypically extreme F2 mice as well as to examine whether the phenotypic effects were found in male and female mice combined or whether there was evidence for gender-specific effects. Statistical significance was reached when $p \leq 0.05$.

Results

Histology of Normal and Fluorotic Enamel

Delayed removal or persistence of matrix proteins was present in the maturing enamel of A/J mice. Hematoxylin and eosin serial cross-sections through the mandibles of A/J mice at the level of the mesial root of the first molar reveal the presence of eosinophilic staining material in the enamel zone following EDTA demineralization in both control (0 ppm F) and treated (50 ppm F) animals (fig. 2a, b). There was greater abundance in the enamel zone following F treatment. The 129P3/J mice had an enamel zone that was barren of material with only a small accumulation of proteinaceous material following F treatment (fig. 2c, d). In order to better understand the composition of the accumulated matrix proteins in fluorotic A/J enamel, we performed immunohistochemistry using anti-amelogenin antibodies. Strong immunoreactivity was found in the enamel zone (fig. 3) and reflected the enamel rod pattern.

Quantitative Trait Loci Identification and Possible Gender Effects

An F2 panel of 458 mice derived from A/J and 129P3/J parental strains were treated with 50 ppm F in their drinking water for 60 days. Monitoring of F exposure, performed by serum F analyses in randomly selected F2 animals representing all 4 TF scales, demonstrated a mean serum F of 12.366 $\pm$ 1.713 $\mu$M. The serum F values obtained in a cross-section of the F2 panel did not significantly differ from serum F determined in comparably treated parental mice (11.296 $\pm$ 3.984 $\mu$M). DF severity in the mandibular incisors was assessed in each individual F2 animal using clinical criteria (modified TF scale; table 1). To maximize the power to detect quantitative trait loci (QTL) contributing to the variation in response to DF, only the phenotypic extreme F2 animals (102

Table 1. Classification of the characteristic clinical appearance of fluorotic incisor enamel in mice as modified from Thylstrup and Fejerskov [1978]

<table>
<thead>
<tr>
<th>TF score</th>
<th>Description</th>
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<tr>
<td>0</td>
<td>The normal translucency of the glossy creamy yellow enamel remains after wiping and drying of the tooth surface.</td>
</tr>
<tr>
<td>1</td>
<td>Thin white opaque lines are seen running across the tooth surface. Such lines are found on all parts of the surface. The lines correspond to the position of the perikymata.</td>
</tr>
<tr>
<td>2</td>
<td>The opaque white lines are more pronounced and frequently merge to form small cloudy areas scattered over the whole surface.</td>
</tr>
<tr>
<td>3</td>
<td>Merging of the white lines occurs, and cloudy areas of opacity spread over many parts of the surface. In between the cloudy areas, white lines can also be seen.</td>
</tr>
<tr>
<td>4</td>
<td>The entire surface exhibits a marked opacity or appears chalky white. Parts of the surface exposed to attrition or wear may appear to be less affected.</td>
</tr>
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From Thylstrup and Fejerskov [1978] and Fejerskov et al. [1994].

Genotyping

Genomic DNA were prepared using the Puregene Tissue kit (Gentra Systems, Minneapolis, Minn., USA) and quantitated by picogreen fluorometry. Genotyping was performed on the Illumina platform at the Mutation Mapping and Developmental Analysis Project (Brigham and Women’s Hospital, Harvard Medical School) [Moran et al., 2006]. Each mouse was genotyped for 354 single nucleotide polymorphism (SNP)-based markers distributed throughout the mouse genome.

Histology

Prior to euthanasia, mice were perfused with 10% neutral buffered formalin (NBF) while under anesthesia. Tissues were collected and further fixed in ice-cold NBF overnight followed by demineralization in 0.2 M EDTA pH 7.2 containing 2% NBF. Mandibles were processed for routine histology. In order to obtain a representative stage of amelogenesis in maturing enamel, cross-sections at the level of the mesial root of the first mandibular molar were obtained and stained with hematoxylin and eosin [Smith and Nanci, 1989]. Serial sections from the same stage of amelogenesis were dewaxed and rehydrated for immunohistochemistry. Following preincubation in 3% hydrogen peroxide in methanol and rinses in 0.03% Tween, sections were subjected to primary polyclonal antibodies (rabbit anti-porcine amelogenin; gift from Dr. James Simmer, University of Michigan, Ann Arbor, Mich., USA) or normal serum as a control. This primary antibody has been shown to have cross-reactivity to rodent amelogenins [Uchida et al., 1991]. Anti-amelogenin immunoreactivity was visualized using the Vectastain Elite ABC kit (Rabbit IgG; Vector Laboratories, Burlingame, Michigan, Ann Arbor, Mich., USA) or normal serum as a control. This primary antibody has been shown to have cross-reactivity to rodent amelogenins [Uchida et al., 1991].
Fig. 2. Hematoxylin and eosin-stained cross-sections of demineralized formalin-fixed paraffin-embedded mandibular incisors at the level of the molar mesial root in A/J (a, b) and 129P3/J (c, d) parental strains treated with 50 ppm F (b, d) or 0 ppm F (a, c). E = Enamel zone; D = dentin.

Fig. 3. Anti-amelogenin immunohistochemical staining of the mandibular incisor enamel zone from an A/J mouse treated with 50 ppm F. Scale bar = 25 μm.
Fig. 4. Genome-wide scan for DF phenotypes for mouse chromosomes 1–19 (excluding X and Y chromosomes). The p values obtained from $\chi^2$ analyses were converted to $-\log$ of p values and plotted on the y axis versus the relative location along each chromosome on the x axis. Data are from 107 mice representing the phenotypic extremes of animals scored as either a clinical DF of 1 or 4 from a panel of 458 F2 mice.

animals with a score of 1 or 4) were genotyped for 354 SNP-based markers distributed throughout the mouse genome. These mice were composed of 55 males and 52 females. $\chi^2$ analysis was performed to compare the genotypic distribution in the 2 groups of phenotypically extreme F2 mice. Significant evidence of association was observed on chromosomes 2 and 11 for a series of consecutive markers ($p < 0.0001$; fig. 4). On chromosome 2, a QTL interval of approximately 77.62 Mb flanked by markers rs13476589 and UT_2_156.443943 was identified, and on chromosome 11, a QTL interval of approximately 74.29 Mb flanked by markers rs3708339 and rs6161623 was identified.

We observed that female mice represented 74% of the F2 animals with a TF score of 1 and male mice represented 76% of the F2 animals with a TF score of 4. No such gender bias was previously observed with the 2 parental strains or with their F1 hybrid progeny (data not shown). These mice were subjected to analyses to examine whether the phenotypic effects were found in both male and female mice and whether there was evidence for gender-specific effects. Significant gender differences in the frequencies of mild (TF = 1) to severe (TF = 4) DF in the F2 panel were observed ($\chi^2$, $p < 0.0001$). Analyses were performed to examine whether the phenotypic gender-specific effects contributed to possible differences in QTL detection. Analyses performed using the markers on chromosomes 2 and 11 which were significant in the mixed-gender mice were also significant when analyses were limited to only the male or female mice. Therefore, the QTL in these 2 regions which influence the variation in response to DF has its effect in mice of both genders. Finally, preliminary analyses suggest that QTL in both chromosomes appear to have an additive effect.

Discussion

The objective of the research was to determine what part of the mouse genome carries susceptibility to fluorosis and to investigate further the nature of the enamel defect in the F-susceptible mice. For the former, F-sensitive and F-resistant inbred mice were mated, generating offspring (F2) that were each unique with respect to the mix of chromosomes they inherited from their parents. Their susceptibility to fluorosis was determined by providing them high F in the drinking water and then staging their fluorosis phenotypes using a modified TF index. As expected, the F2 mice varied with respect to their susceptibility to fluorosis, but unexpectedly, F2 males and females varied in their susceptibility. To characterize what parts of the genome of these mice were inherited from the F-susceptible and F-resistant parents, each F2 mouse was genotyped, that is, characterized for the presence of 354 SNPs distributed throughout the genome. Statistical analyses were performed for the F2 mice that were the most F-resistant and F-susceptible to determine which parts of the genome correlate with these phenotypes. Our investigations of DF susceptibil-
ity/resistance QTL using A/J and 129P3/J inbred mice in the generation of F2 progeny was capable of detecting QTL on chromosomes 2 and 11. The QTL intervals detected in this study contain nearly 2,000 genes. More importantly, lack of significant evidence of association observed on chromosomes X, 3, 5, 7 or 9 suggests a small role of amelogenin, ameloblastin, enamelin, amelotin, Klk-4 or Mmp20 in DF susceptibility/resistance in our animal model. We have no ready explanation for the gender differences observed in our F2 panel. Sex or gender differences in DF affecting humans have not been observed. Future studies will focus on narrowing the QTL intervals detected and identifying candidate genes for further interrogation.

Separately, the nature of the enamel defect in the F-susceptible mouse was investigated. Histological analyses of the maturing EDTA-soluble enamel from the F-susceptible A/J mouse strain following F exposure demonstrated an accumulation of matrix proteins composed in part of amelogenins. Interestingly, some retention of matrix proteins was present in the 0 ppm F treatment group of the A/J strain. The enamel zone of comparably staged 129P3/J incisor remained devoid of any proteinaceous material. The accumulation of enamel matrix proteins in the fluorotic enamel found in the A/J mice is consistent with many previous studies [Den Besten, 1986; Den Besten and Thariani, 1992; Den Besten, 1999]. We found detectable accumulation of matrix proteins in the nonfluorosed enamel of A/J mice suggesting that inherent differences in tooth enamel formation may exist between these 2 strains of mice. Previously, we observed baseline differences in enamel quality between A/J and 129P3/J mice. Using microhardness as a criteria, the incisor enamel of 129P3/J mice is approximately 22% harder than that of A/J [Vieira et al., 2005]. More notably, we found that F treatment (50 ppm F) resulted in a greater loss of enamel microhardness in the A/J strain (71% reduction) compared to a 15% reduction in the 129P3/J strain. While the retention/persistence of matrix proteins in maturing enamel of A/J mice was greatly enhanced by systemic F exposure and likely contributed directly to the DF phenotype present in this strain, new questions emerge regarding genetically determined variability in normal enamel formation.

References


