

Benign Migratory Glossitis: A Review

A Hooda, M Rathee, J Gulia, S Yadav

Citation

A Hooda, M Rathee, J Gulia, S Yadav. *Benign Migratory Glossitis: A Review*. The Internet Journal of Family Practice. 2009 Volume 9 Number 2.

Abstract

Benign migratory glossitis also known as geographical tongue is a recurrent condition of unknown etiology characterized by loss of epithelium particularly of the filiform papillae on the dorsum of the tongue. Clinically the appearance is of multifocal, circinate, irregular erythematous patches bounded by slightly elevated, white colored keratotic bands. Several etiologic factors have been proposed. The condition may remain asymptomatic or patients may present with complains of pain and burning sensation. A review of the current opinion on benign migratory glossitis is presented.

INTRODUCTION

Geographic tongue is a transient and recurrent condition characterized by periodic localized loss of epithelium particularly of the filiform papillae on the dorsum of the tongue. The pattern of the red map like areas with white borders which occasionally migrate across the tongue gives an abnormal appearance resembling a map and may involve buccal mucosa.¹ Several factors proposed as possible causative agents of geographic tongue, which include local factors, hormonal disturbances and various systemic diseases. Geographic tongue has a wide spectrum of appearance and variable symptomatology. The clinical presentation may vary from asymptomatic to a painful and burning ulceration. Management of geographic tongue depends upon the clinical presentation and the underlying etiology. A review of the current concept in causes and management is being presented.

REVIEW

Benign migratory glossitis or geographic tongue is common benign disorder of unknown etiology. The epithelium of the tongue is affected with loss of filiform papillae leading to smooth ulcer like lesions that rapidly change the color and size. The lesions commonly occur on the tip, lateral borders, dorsum of the tongue and sometimes extend to the ventral portion of the tongue.² The prevalence rate is between 1.0% and 2.5%.³ According to Jainkittivong and Langlais⁴ the highest incidence of geographic tongue is in the 20-29 age group. Shulman and Carpenter⁵ on the other hand found no relation between benign migratory glossitis and age among US adults. A higher female preponderance is reported.

Jainkittivong and Langlais observed higher rates in females (1.5:1) between ages 9 and 79 in a population in Thailand⁴. Similarly Marks and Simons reported a female to male ratio of 1.4:1 in the age group of 3 to 77 in an Australian population.⁶ Some authors on the other hand found that gender was not important to the incidence of geographic tongue.⁷

It is reported to begin in childhood and is most frequently observed in children four to four and half years of age.⁸ It is localized most commonly on the dorsum of the tongue, lateral borders and the tip of the tongue. It is characterized by discrete smooth reddened areas, usually slightly raised with pale yellow or white borders. When observed over a period of hours or days the denuded patches may change drastically in size and shape, often appearing to migrate across the surface of the tongue or disappearing for widely varying period of time. The pattern has been likened to land masses and oceans on a map, from which the synonym geographic tongue was derived.^{9,10}

Hashemipour et al in a study of 837 students found that 104 students had geographic tongue. Most (74%) lesions manifested a typical appearance consisting of a central atrophic area bounded by a raised white circinate line and the remaining lesions were characterized as solely atrophic patches. The most common site of the lesions was the lateral borders of the tongue.¹¹

A process similar to geographic tongue when occurs in other areas of oral mucosa is called "ectopic geographic tongue". This was first described by Cooke (1955) under the name of

erythema migrans.¹² In the literature several other names are also in use for this condition namely: geographic stomatitis^{13,14}, stomatitis areata migrans¹⁵, erythema migrans¹⁶ and migratory stomatitis¹⁷. It is emphasized that the ectopic geographic tongue is the same process as the geographic tongue involving other areas of the oral mucosa.¹⁸

The etiology and pathogenesis remains obscure. Many risk factors have been proposed including hormonal disturbances¹⁹, oral contraceptive use²⁰, juvenile diabetes mellitus²¹, pustular psoriasis^{22,23}, allergic conditions such as atopy, hay fever and rhinitis^{24,25}, fissured tongue^{26,27}, Robinow's syndrome²⁸, Reiter's syndrome²⁹, Down syndrome^{30,31}, psychological factors³², nutritional deficiencies³³, lithium therapy^{23,34}, familial predisposition³⁵⁻³⁷, Fetal hydantoin syndrome³⁸ and Aarskog's syndrome³⁹.

HEREDITY

A tendency for familial occurrence of this condition has been suggested. Redman et al reported a higher prevalence of benign migratory glossitis among first degree relatives of affected University of Minnesota students than among those of control students as well as among population from which the students were drawn. They suggested a polygenic model of hereditary transmission, with a threshold for susceptibility to environmental factors.¹⁰ Eidelman et al also suggested that geographical tongue is an inherited condition with a polygenic mode of condition.²⁷

ASSOCIATION WITH PSORIASIS

It has been proposed that benign migratory glossitis is an isolated oral manifestation of psoriasis rather than a mere association.⁴⁰ The relationship of human leukocyte antigens (HLA) with psoriasis has been extensively investigated. Associations of HLA- Cw6 presents a particularly strong association irrespective of the racial or ethnic group suggesting that Cw6 itself or closely linked gene in strong linkage disequilibrium, is the major HLA-linked susceptibility gene for psoriasis.⁴¹ Wysocki and Daley reported the HLA phenotypes of seven patients with benign migratory glossitis and juvenile diabetes and found that none of the cases had Cw6 antigen and five had DR3 or DR4 which are the classical diabetes association²¹. Marks and Tait in a study of ninety five patients with benign migratory glossitis found a marginal increase in the frequency of B15.⁴² Fenerly et al in a study of 50 Greek patients showed increased frequency of DR5 and DR6 and decreased frequency of B51³⁵. Gonzaga et al in a study of 22 patients reported a highly significant association of Cw6 with both

psoriasis and benign migratory glossitis, with this antigen being present in 59.1% of the patients of psoriasis, 43.8% of the patients with benign migratory glossitis and in only 12.6% of the controls⁴¹. These reports reinforce the concept of a pathogenetic relationship between migratory glossitis and psoriasis.

Further histological findings and the parallel improvement of psoriatic skin lesion and tongue lesion with systemic retinoid treatment supports this hypothesis. The infiltrate in oral psoriatic lesions consists mainly of T cells and macrophages, particularly CD4 positive cells and immune-histochemical studies of geographic tongue show similar abundance of CD4 positive cells in the sub-epithelial cell infiltrate⁴³. However other researchers have reported that geographic tongue is uncommon in patients with psoriasis⁴⁴.

ALLERGY

Patients with personal or family history of asthma, eczema and hay fever or elevated total serum immunoglobulin E levels may be more likely to have a geographic tongue. Psychosomatic factors, which probably contribute to both geographic tongue and atopy, may explain the high prevalence of this disorder in atopic patients. Goregen et al recently (2010) used patch test and prick tests to test different mechanisms associated with allergy. The skin prick test measures specific IgE anti bodies in the serum and is used to indicate sensitization. The patch test is helpful to determine delayed-type allergic reactions. The authors reported that performing both tests in combination improves the diagnostic efficacy of predisposition of allergy patients with benign migratory glossitis.⁴⁵

Marks and Simons found a significantly increased frequency of atopy among patients with geographic tongue as compared to normal population. The prevalence of the HLA antigen B15 was found to be significantly elevated in cases with geographic tongue when compared to a normal population⁶. McLendon and Jaeger reported children with milk allergy and stated that benign migratory glossitis occurred in a significant proportion of these patients.⁴⁶

HORMONAL FACTORS

It has been reported that hormonal fluctuations can affect the geographical tongue. The phase of the oral contraceptives cycle affected the initiation and duration of circinate lesions in women with geographic tongue; the changes were most severe on the day seventeenth of the cycle.²⁰ Further about 8% of the patients with juvenile diabetes mellitus have a

3 of 7

Figure 1

Fig 1: Showing the geographic tongue with multiple lesions involving the anterior two third of the tongue. The erythematous atrophic areas are seen.



Multiple lesions are commonly seen.^{1,4} The white margin is the earliest clinical change with progression to atrophic erythematous areas being a later stage before healing. (Fig2).

Figure 2

Fig 2: Showing the white margins of the lesion on the left side of the tongue surrounding an erythematous atrophic area



The erythema represents atrophic filiform papillae with loss of keratin, slightly depressed atrophic areas are surrounded with a sharply defined edge and slightly elevated white borders and degranulation of the overlying mucosa. The morphology is changing and it thus gives the appearance of migrating across the tongue.^{1,10,11}

Similar appearing lesions are occasionally seen in the other areas of oral mucosa such as soft palate, buccal mucosa, floor of mouth, gingiva, and uvula. These are referred to as geographic stomatitis and these conditions represent the non-lingual counterpart of geographic tongue.¹⁴⁻¹⁸

HISTOPATHOLOGY

Histopathologic examination may reveal an acute and chronic inflammatory infiltrate in the submucosa with epithelial edema. It is associated with neutrophils forming microabscess. In the sections of areas corresponding to the red patches increased edema is noted in the acanthotic epithelium and differentiation into filiform papillae is lacking. The inflammatory cells accumulate in the uppermost layers of the epithelium and form micro-abscess.

The areas from middle of the red patch show complete absence of the filiform papillae and parakeratotic layers. The white elevated areas of the lesions include subepithelial neutrophil infiltrates and micro-abscesses, leukocyte invasion into the epithelial layer, intraepithelial edema, rupture of cell junction, glycogen deposits in the epithelial cells and exfoliation of necrotic cells in the surface layer.^{1,52} Scanning electron microscopy has revealed that the surface of the geographic tongue contains two types of abnormal mucosa: an atrophic area on which the hairs of filiform papillae are absent but the bodies appear typical, and a white margin of desquamating cells. Micro-fissures are located between atrophic and normal mucosa.⁵³

DIAGNOSIS

Diagnosis is based upon history and clinical examination. Routine laboratory tests are usually normal. Biopsy and histological examination of the lesions is usually not required considering the benign nature of the disease but may assist in reassuring patients more so with cancer phobia of the benign nature of the disease.^{2,33}

DIFFERENTIAL DIAGNOSIS

Differential diagnosis includes candidiasis, psoriasis, Reiter's syndrome, leukoplakia, lichen planus, systemic lupus erythematosus, herpes simplex, and drug reaction. In children local trauma and severe neutropenia should be excluded.² Candidiasis is the most common oral fungal infection in humans. Leukoplakia the most common premalignant lesion of oral mucosa is defined as a white patch or plaque that cannot be removed by vigorous scraping and cannot be classified as specific disease entity. The patients main concern in geographical tongue is that his/ her lesions could represent cancer leukoplakia should be considered a differential diagnosis.^{54,55}

TREATMENT

Patients do not usually require treatment apart from reassurance. Various symptomatic treatments have been tried and include fluids, acetaminophen, mouth rinsing with topical anesthetic agent, antihistaminics, anxiolytics and steroids.² Helfman reported satisfactory results after treating three patients with topical tretinoin. Vitamin A therapy resulted in partial improvement in some patients.⁵⁶ The topical factors that exacerbate patient's symptoms such as very hot, spicy or acidic food, and dried salty nuts should be avoided.² Abe et al reported marked improvement in a 54 year-old female with persistent and painful benign migratory glossitis (BMG), for about five years by systemic

administration of cyclosporin. The systemic treatment of cyclosporin microemulsion pre-concentrate, 3 mg/kg/day, resulted in a satisfactory improvement. Two months later, patient was started on maintenance therapy with cyclosporin microemulsion pre-concentrate dosage to 1.5 mg/kg/day.⁵⁷

CONCLUSION

Benign migratory glossitis or geographic tongue is common benign disorder of unknown etiology. The clinical presentation may vary from asymptomatic to a painful and burning ulceration. Management of geographic tongue depends upon the clinical presentation, the underlying etiology and should include reassuring the patients more so with cancer phobia about the benign nature of the disease.

References

1. Marks R, Radden BG. Geographic tongue: a clinicopathology review. *Aust J Dermatol* 1981; 22: 75-9.
2. Assimakopoulos D, Patrikakos G, Fotika C, Elisaf M. Benign migratory glossitis or geographic tongue: an enigmatic oral lesion. *Am J Med* 2002; 113:751-5.
3. Kovac-Kovacic M, Skaleric U. The prevalence of oral mucosal lesions in a population in Ljubljana, Slovenia. *J Oral Pathol Med* 2000; 29:331-5.
4. Jainkittivong A, Langlais RP. Geographic tongue: clinical characteristics of 188 cases. *J Contemp Dent Pract* 2005; 1:123-35.
5. Shulman JD, Carpenter WM. Prevalence and risk factors associated with geographic tongue among US adults. *Oral Dis* 2006; 12:381-6.
6. Marks R, Simons MJ. Geographic tongue—a manifestation of atopy. *Br J Dermatol* 1979; 101:159-62.
7. Avcu N, Kanli A. The prevalence of tongue lesions in 5150 Turkish dental outpatients. *Oral Dis* 2003; 9:188-95.
8. Rioboo-Crespo Mdel R, Planells-del Pozo P, Rioboo-Garcia R. Epidemiology of the most common oral mucosal diseases in children. *Med Oral Patol Oral Cir Bucal* 2005; 10:376-87.
9. Miloglu O, Göregen M, Akgül HM, Acemoglu H. The prevalence and risk factors associated with benign migratory glossitis lesions in 7619 Turkish dental outpatients. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2009; 107:e29-33.
10. Redman RS, Shapiro BL, Gorlin RJ. Hereditary component in the etiology of benign migratory glossitis. *Am J Hum Genet* 1972; 24: 124-33.
11. Hashemipour M, Rad M, Dastboos A. Frequency, clinical characteristics and factors associated with geographic tongue. *Shiraz Univ Dent J*. 2008; 9: 83-92.
12. Cooke BED. Erythema migrans affecting the oral mucosa. *Oral Surg Oral Med Oral Pathol* 1955; 8: 164-7.
13. Hume WJ. Geographic stomatitis: a critical review. *J Dent* 1975; 3:25-43.
14. Donelli RA. Geographic stomatitis (tongue and mucosae). *J Clin Stomatol Conf*. 1964; 5:21-22.
15. Saprio SM, Shklar G. Stomatitis areata migrans. *Oral Surg Oral Med Oral Pathol* 1973; 36: 28-33.
16. Rood JP. An unusual presentation of erythema migrans. *J Dent* 1974; 2: 207-8.
17. Zingale J. Migratory stomatitis. *J Periodontol* 1977; 48: 298-302.

18. Grinspan D, Blanco GF, Aguero S, Bianchi O, Stringa S. Ectopic Geographic tongue and AIDS. *Int J Dermatol* 1990; 29: 113-6.
19. Kullaa-Mikkonen A, Kotilainen R, Alakuijala P. Sialochemistry in mucosal lesions of the tongue: electrolytes and total protein. *Int J Oral Maxillofac Surg* 1986; 15:318-21.
20. Waltimo J. Geographic tongue during a year of oral contraceptive cycles. *Br Dent J* 1991; 171:94-6.
21. Wysocki GP, Daley TD. Benign migratory glossitis in patients with juvenile diabetes. *Oral Surg Oral Med Oral Pathol* 1987; 63:68-70.
22. Zhu JF, Kaminski MJ, Pulitzer DR. Psoriasis: pathophysiology and oral manifestations. *Oral Dis* 1996; 2:135-44.
23. Zargari O. The prevalence and significance of fissured tongue and geographical tongue in psoriatic patients. *Clin Dermatol* 2006; 31:192-5.
24. Marks R, Scaff CE, Yap V, Verlinden V, Jolley D, Champell J. Fungiform papillary glossitis: disease in the mouth? *Br J Dermatol* 2005; 153:740-5.
25. Marks R, Czarny D. Geographic tongue: sensitivity to the environment. *Oral Surg Oral Med Oral Pathol* 1984; 58:156-9.
26. Yarom N, Cantony U, Gorsky M. Prevalence of fissured tongue geographic tongue and median rhomboid glossitis among Israeli adults of different ethnic origins. *Dermatol* 2004; 209:88-94.
27. Eidelman E, Chosack A, Cohen T. Scrotal tongue and geographic tongue: polygenic and associated traits. *Oral Surg Oral Med Oral Pathol* 1976; 42:591-6.
28. Cerqueira DF, de Souza IP. Orofacial manifestations of a patient with Robinow's syndrome: a case report in pediatric patient. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2008; 105:353-7.
29. Fotiou G, Laskaris G. Reiter's syndrome oral manifestations. *Hell Stomatol Chron* 1988; 32: 148-51.
30. Daneshpazhooh M, Nazemi TM, Bigdeloo L, Yoosefi M. Mucocutaneous findings in 100 children with Down syndrome. *Pediatr Dermatol* 2007; 24:317-20.
31. Ercis M, Balcı S, Atakan N. Dermatological manifestations of 71 children admitted to clinical genetics unit. *Clin Genet* 1996; 50:317-20.
32. Redman RS, Vance FL, Gorlin RJ, Peagler FD, Meskin LH. Psychological component in the etiology of geographic tongue. *J Dent Res* 1966; 4:1403-8.
33. Banoczy J, Szabo L, Csiba A. Migratory glossitis. A clinical- histological review of seventy cases. *Oral Surg Oral Med Oral Pathol* 1975; 39:113-21.
34. Patki AH. Geographic tongue developing in a patient on Lithium Carbonate therapy. *Int J Dermatol* 1992; 31:368-9.
35. Fenerli A, Papanicolaou S, Papanicolaou M, Laskaris G. Histo compatibility antigens and geographic tongue. *Oral Surg Oral Med Oral Pathol* 1993; 76:476-9.
36. Pavelic J, Gall-Troselj K, Mravak-Stipetic M, Pavelic K. The p53 and nm23-H1 genes are not deleted in oral benign epithelial lesions. *Anticancer Res* 1998; 18:3527-31.
37. Eidelman E, Chosack A, Cohen T. Scrotal tongue and geographic tongue: polygenic and associated traits. *Oral Surg Oral Med Oral Pathol* 1976; 42:591-6.
38. Nanda A, Kaur S, Bhakoo ON, Kapoor MN, Kanwar AJ. Fetal hydatoin syndrome: a case report. *Pediatr Dermatol* 1989; 6:130-3.
39. Hassinger DD, Mulvihill JJ, Chandler JB. Aarskog's syndrome with Hirschprung's disease, midgut malrotation and dental anomalies. *J Med Genet* 1980; 17: 235-8.
40. Pogrel MA, Cram D. Intraoral findings in patients with psoriasis with a special reference to ectopic geographic tongue (erythema circinata). *Oral Surg Oral Med Oral Pathol* 1988; 66:184-9.
41. Gonzaga HFS, Torres EA, Alchorne MMA, Gerbase-Delima M. Both psoriasis and benign migratory glossitis are associated with HLA-Cw6. *Brit J Dermatol* 1996; 135:368-70.
42. Marks R, Tait B. HLA antigens in geographic tongue. *Tissue Antigens* 1980; 15:60-2.
43. Espelid M, Bang G, Johannessen AC, Leira JL, Christensen O. Geographic stomatitis: report of 6 cases. *J Oral Pathol Med* 1991; 20: 425-8.
44. Hietanen J, Salo OP, Kanerva I, Juvakoski T. Study of the oral mucosa in 200 consecutive patients with psoriasis. *Scand J Dent Res* 1984; 92: 50-4.
45. Goregen M, Melikoglu M, Miloglu O, Erdem T. Predisposition of allergy in patients with benign migratory glossitis. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2010; 110: 470-4.
46. McLendon PA, Jaeger DS. Milk intolerance, the cause of a nutritional entity: a clinical study. *South Med J* 1943; 36: 571-5.
47. Chosack LJ, Zadik D, Eidelman E. The prevalence of scrotal tongue and geographical tongue in 70359 Israeli schoolchildren. *Community Dent Oral Epidemiol* 1974; 2: 253-7.
48. Wright V, Reed WB. The link between Reiter's syndrome and psoriatic arthritis. *Ann Rheum Dis* 1964; 23: 12-21.
49. Kulka JP. The lesions of Reiter's syndrome. *Arthritis Rheum* 1962; 5:195-201.
50. Weathers DR, Baker G, Archard HO, Brukes EJ Jr. Psoriasiform lesions of the oral mucosa (with emphasis on "ectopic geographic tongue"). *Oral Surg Oral Med Oral Pathol* 1974; 37: 872-88.
51. Sigal MJ, Mock D. Symptomatic benign migratory glossitis: report of two cases and literature review. *Pediatr Dent* 1992; 14:392-6.
52. Plackova A, Skach M. The ultrastructure of geographic tongue. *Oral Surg Oral Med Oral Pathol* 1975; 40: 760-8.
53. Kullaa-Mikkonen A. Geographic tongue: an SEM study. *J Cutan Pathol* 1986; 13: 154-62.
54. Binmadi NO, Jham BC, Meiller TF, Scheper MA. A case of deeply fissured tongue. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2010; 109: 659-63.
55. Greer RO. Pathology of malignant and pre-malignant oral epithelial lesions. *Otolaryngol Clin North Am* 2006; 39: 249-75.
56. Helfman RJ. The treatment of geographic tongue with topical retin-A solution. *Cutis* 1975; 50: 41-6.
57. Abe M, Sogabe Y, Syuto T, Ishibuchi H, Yokoyama Y, Ishikawa O. Successful treatment with cyclosporine administration for persistent benign migratory glossitis. *J Dermatol* 2007; 34: 340-3.

Author Information

Anita Hooda, MDS Prosthodontics

Department of Oral Anatomy, Pt. B.D Sharma University of Health Sciences

Manu Rathee, MDS Prosthodontics

Assistant Professor, Department of Prosthodontics, Pt. B.D Sharma University of Health Sciences

Joginder Singh Gulia, MS Otorhinolaryngology

Professor, Department of Otorhinolaryngology, Pt. B.D Sharma University of Health Sciences

Samar Pal Singh Yadav, MS Otorhinolaryngology

Senior Professor, Department of Otorhinolaryngology, Pt. B.D Sharma University of Health Sciences