

Case report

Dental considerations in a 4-year-old girl with Lennox-Gastaut Syndrome. Case report and literature review.

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Abstract

We present the developmental, oral, clinical, radiographic findings and oral treatment of a 4-year-old girl **presenting with** Lennox-Gastaut syndrome (LGS), which is a severe disabling childhood epilepsy diseases that is treated with one or multiple anti-epileptic drugs (AEDs). The

25 | child was wheel-chair bound, developmentally delayed, gastrostomy tube (G-tube) fed, and
26 | suffered from multiple seizures and infantile spasms.⁵ The child's medical history included an
27 | under-developed pituitary gland, gastro esophageal reflux disease, vision and hearing
28 | impairment, history of chronic aspiration pneumonia, and allergies. The oral findings included
29 | no carious lesions, heavy calculus accumulation, spontaneous bleeding from the gingiva,
30 | generalized gingival hyperplasia (GH) and abnormal increased mobility in several
31 | primarydeciduous teeth. The comprehensive radiographic and clinical examination and the
32 | treatment under general anesthesia are described. ~~The etiologies of the calculus accumulation~~
33 | ~~and GH are reviewed.~~

34 | Key words: calculus, gingival overgrowth, anti-seizure medication, Lennox-Gastaut Syndrome

36 | Introduction

37 | Lennox-Gastaut syndrome (LGS) is a severe and disabling childhood epilepsy that is
38 | characterized by a triad of symptoms: 1) ~~generalized treat~~ seizures resistant to multiple therapies;
39 | 2) slowness of intellectual growth and cognitive impairment; 3) a specific electroencephalogram
40 | (EEG) disturbance called a slow spike-and-wave pattern that is present when the child is
41 | awake.¹⁻⁵ LGS patients may have multiple daily seizures that may cause sudden and
42 | unpredictable stiffening followed by a drop to the ground; this being a key diagnostic feature.⁵⁻⁷
43 | The pharmacologic treatment may include one or multiple antiepileptic drugs (AEDs),⁴ some of
44 | which have the potential to induce gingival hyperplasia (GH).

45 | A review of the literature identified only one report of the oral findings in a LGS patient, ~~of a~~
46 | 26-year-old female who had macroglossia, supragingival as well as subgingival calculus, red,

47 swollen and friable gingiva with generalized bleeding and localized suppuration, and gingival
48 recession.⁴ The present manuscript includes an additional , comprehensive case report of a 4-
49 year-old girl with LGS, and presents a review of the literature on LGS and related anti-seizure
50 medication that may have-induced gingival overgrowth.

51

52 Case presentation

53 A 4.5-year-old Caucasian female with LGS was referred to a University Clinic for dental
54 treatment. The medical history indicated that she was born at 32 weeks of gestation, along with
55 her healthy twin. The patient had infantile seizures and spasms 15-16 times per day and was
56 diagnosed with LGS. As a result of her condition, she experienced Her medical history was
57 significant for developmental delay, wheelchair-bound, had a gastrostomy tube (G-tube), under-
58 developed pituitary gland, gastro esophageal reflux disease, vision and hearing impairments,
59 history of chronic aspiration pneumonia, allergies to Depakote and Amoxicillin and
60 leukodystrophy (degeneration of the white matter in the brain⁸). Her medications included;
61 Vigabatrin, Clobazam, Topiramate, Fycompa, Diazepam and Rufinamide reducing the daily
62 seizures to 3-6, and Albuterol/atropine via nebulizer. At the time of this study, she had recently
63 been hospitalized due to seizures,~~Recent hospitalizations resulting from seizures,~~ chronic
64 pneumonia, and adrenocorticotrophic hormone therapy. Her~~The~~ surgical history included adenoid
65 and tonsils removal, Nissen fundoplication with hernia repair, and G-tube placement. The chief
66 complaint as expressed by her mother was risk of aspirating exfoliating primary teeth: the
67 previous night the patient had a seizure, after which she was “choking and was missing a lower
68 tooth that was swallowed or aspirated”.

69 On examination, she had no apparent respiratory difficulties, but was non-verbal, had a small
70 “hypoplastic” face, inability to cooperate, extensive drooling, short stature and slight
71 overweight.⁹ A limited oral examination revealed sialorrhea, deciduous primary dentition with
72 missing mandibular deciduous primary central incisors, heavy calculus on the majority of teeth
73 surfaces, abnormal mobility (2-3 mm) in both mandibular primary deciduous lateral incisors
74 (teeth #N and #Q), as well as generalized moderate GH. Tongue size appeared normal. A chest
75 radiograph did not reveal tooth aspiration.

76 The patient MG was admitted to the hospital the day before the dental treatment under
77 general anesthesia (GA), maintained with intravenous fluid to avoid the conflict between being
78 fed nil per os and her need for frequent gastrostomy tube (G-tube) feeding. The mother reported
79 that the patient was apparently having pain while grinding her teeth. Under GA, a-radiographic
80 and clinical examinations revealed no caries, no evidence of dental pulp pathology (Figure 1),
81 all maxillary primary deciduous incisors (Teeth D, E, F, G), and both mandibular lateral
82 primary deciduous incisors (Teeth N and Q) had abnormal mobility (about 3 mm), nearly all
83 teeth were covered with heavy calculus (Figure 2), generalized moderate GH, and a band of
84 gingiva over the occlusal surface of the mandibular right first primary deciduous molar (Teeth S,
85 Figure 3A Figures 2,3, 4A), and gingiva over the occlusal surface of the maxillary right first
86 deciduous primary molar (Tooth B, Figure 4a). The GH was non-hemorrhagic, soft, slightly
87 fluctuant and pink (Figures 2, 3a&b, 4a&b). Calculus removal was accomplished with an
88 ultrasonic and hand instrumentation, followed by an application of a fluoride varnish. The
89 gingival tissue over teeth B and S were removed with a surgical blade (Figures 3b & 4b). Teeth #
90 D, E, F, G, N and Q were extracted. The post-operative recovery was uneventful .The patient’s
91 irritability associated with her oral pain has subsided significantly.-

92 Discussion and literature review

93 | Dr. William Lennox, first described LGS in the 1930s, Lennox and Davis later reported its
 94 | triad, which was further expanded by Gastaut.¹¹⁻¹² The median onset age of LGS is about 4 years
 95 | (range: 0.6-28.9 years) with a peak onset of 5 years.¹³⁻¹⁴ LGS is uncommon (3-10% of childhood
 96 | epilepsy) and has a mortality rate ranging from 3% to 7%.^{2, 3, 12} The tonic seizures are
 97 | characterized by an EEG diffuse high voltage slow wave followed by generalized low voltage
 98 | fast activity, reflecting sustained fast neurological firing over a wide cortical area.^{5, 15} 80% of
 99 | LGS patients will continue to have seizures into adulthood.^{2, 16}

100 | Based on our literature review, this is the second case in which the oral characteristics of
 101 | LGS are described, and the first one in a child. In this case the dental consideration included
 102 | behavioral and management issues, gingival hyperplasia as a result of side effects caused by anti
 103 | seizure medication, poor oral hygiene (OH), and a risk of aspiration from loose teeth and
 104 | difficulties in swallowing. Comparison of both cases is restricted by the patients' different age
 105 | groups; the previous report was in a 26-year-old female.⁴ Both cases received AEDs
 106 | and had GH and severe calculus accumulation; the previous case had periodontitis and
 107 | macroglossia that encumbered proper OH while in the present case the tongue size was normal
 108 | and there was increased abnormal tooth mobility; with no radiographic evidence of alveolar
 109 | bone loss. Oral pain was reported in the previous case associated with gingival swelling, gingival
 110 | recession and periodontitis. In the younger patient while in the present case, pain was assumed
 111 | to be related to biting on the gingival tissue over the occlusal surfaces.

112 | GH commonly starts with the eruption of the permanent dentition and may be influenced by
 113 | genetic predisposition.¹⁷ However, in the present case there was no history of GH in the
 114 | family,¹⁷ indicating that the GH may have been caused by one or more AEDs most likely

115 | Vigabatrin. The aim of AEDs is to control or decrease seizures without producing unacceptable
116 | adverse effects that impair quality of life.¹⁷ However, AEDs have been most frequently
117 | associated with drug side effects ~~adverse drug reactions~~.¹⁷ The pharmacologic treatment of LGS
118 | includes AEDs such as Vigabatrin, Valproates, Felmabate, and Benzodiazepines which may
119 | potentiate each other's side effects, as in cases in which GH is potentiated by the combination of
120 | phenytoin and calcium channel blockers, or cyclosporine and calcium channel blockers.¹⁸⁻²⁰

121 | Interestingly, multiple AEDs have a multiplier n-additive effect on GH, that might explain
122 | the additive effect of multiple anticonvulsant therapy to GH.²⁵

123 | GH might include an abundance of dense connective tissue or acellular collagen that can be
124 | an impediment to tooth eruption.^{36, 37} Delayed eruption has also been associated with severe
125 | bruxism in children with cerebral palsy.^{38, 39} In the present case, the primary dentition was
126 | normal.⁴⁰ However, the clinical crowns of the deciduous primary teeth appeared shorter than
127 | normal and there was gingival tissue at the occlusal surfaces of teeth B and S, suggesting a
128 | combination of GH and delayed eruption that could be related to the GH and bruxism (Figures
129 | ,3a, 3b).

130 | Despite the positive correlation between plaque scores, gingival inflammation, and severity
131 | of GH in children, the role of OH as an etiologic factor for GH has not yet been fully clarified
132 | since most of the studies have been cross-sectional.^{19, 25} However, the relevance of OH is
133 | emphasized in the previously reported LGS case, in which non-surgical periodontal therapy was
134 | effective in controlling periodontal disease, and prevention of oral diseases is preferable for a
135 | high-risk patients.⁴ In the present case however, OH performance is complicated by the child's
136 | inability to perform the most simple measures and to cooperate with her parents.

137 A full mouth gingivectomy in the primary dentition was reported by Breen et al. (2009) in a
138 case of a 28 month old with hereditary gingival fibromatosis in which only 4 mandibular teeth
139 were partially erupted.¹⁷ In the present case, we included the removal of the gingival tissue from
140 the occlusal surfaces of the primary molars that most likely were the origin of oral pain (Figures
141 3b & 4b); in retrospective, a gingivectomy could have been adequate for the maxillary right
142 primary cuspid and lateral incisor that had minimal clinical crowns (Figure 4a). ~~;~~ ~~†~~ The patient
143 will continue to be under follow-up and will be scheduled for gingivectomy if required.

144 Children and adolescents who are unable to meet their nutritional needs orally and depend
145 on ~~GT-tube~~ feeding ~~at~~ are at a significantly increased risk of poor oral health, specially tartar,
146 accumulation an subsequent gingivitis.^{10, 41, 46} Therefore, ~~in~~ the present case, the possibilities of
147 recurrence of calculus accumulation are high. ~~Based on our search of the literature, it appears~~
148 ~~that this is the youngest case reported with severe generalized calculus accumulation.~~

149 Aspiration of exfoliating primarydeciduous teeth is apparently ~~most uncommon or non-~~
150 ~~reported since our review of the literature disclosed only one~~ uncommon. A case of aspiration of
151 a maxillary primary cuspid by a 9 year 11 month old child with cerebral palsy was reported, the
152 authors have ~~emphasized~~ ing the fact that the possibility of aspiration of primarydeciduous teeth
153 is ~~exacerbated~~ in debilitated patients.⁴⁷ Also, avulsion of primary teeth due to trauma and their
154 aspiration is possible.⁴⁸ This emphasizes the need ~~to consider~~ ~~the need~~ to investigate refer
155 children who “lost” a primary tooth that cannot be found ~~to~~ using a chest radiograph, especially in
156 children with developmental disturbances, ~~or~~ and a history of aspiration pneumonia which
157 involves the entry of infectious pharyngeal contents into the lower airway.⁴¹ Relevant is the fact
158 that low salivary flow associated with gastric tube (GT) feeding may predispose the growth of

159 salivary bacteria that, when mixed with food or liquid, provide a substantial inoculum to the
 160 lungs if aspirated.⁴¹

161 In conclusion, LGS in young children presents a significant challenge to the dental professional
 162 including GA consideration, G-tube issues, poor oral hygiene and gingival hyperplasia, both. The
 163 neurologist and the pediatric dentist should be aware of the potential complication and work as team
 164 on behalf of the patient and the family of the LGS patient.

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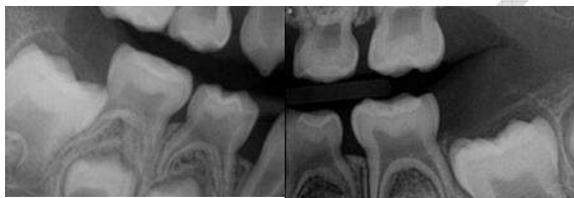
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281 Figure1: a radiographic examinations of the patient achieved during general anesthesia revealed
282 no caries, and no evidence of dental pulp pathology. The lower deciduous incisors are missing

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285 Figure-2

286 Teeth covered with heavy plaque and calculus.

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UNDER PEER REVIEW



290 Figure -3

291 Gingival overgrowth and gingivitis affecting the maxillary R quadrant

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UNDER PEEK



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297 Figure-43a

298 Generalized moderate gingival hyperplasia, and a band of gingiva over the occlusal surface of299 the mandibular right first deciduous molar

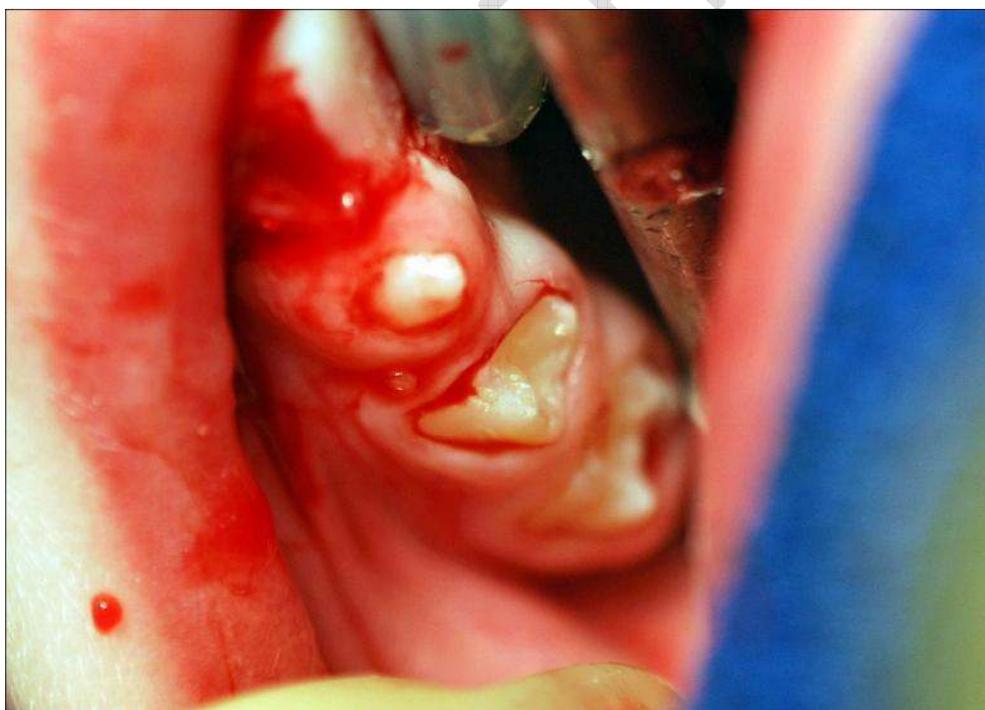
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301 Figure 4b. Gingival tissue over teeth B and S after removal with a surgical blade



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303 | Figure-4a



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305 | Figure-4b

UNDER PEER REVIEW