

Dental Remineralization: Simplified

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Throughout our patients' lives we interact at regular intervals, assessing and treating as their oral condition requires. The ongoing progression of disease in dental hard and soft tissue is often a multifactorial downward spiral. It does not have to be. The dental team must fully understand the disease process and then proactively intervene to slow or preferably stop its progress. This is the concept of **Proactive Intervention Dentistry**. Tools and techniques are needed to provide these services. These tools are readily available and can be easily incorporated into daily practice. This article focuses on **dental hard tissue** and the various products and systems demonstrated to be beneficial in reversing and controlling the caries process. These systems should be used proactively when the patient requires extra help in maintaining hard tissue health.

DEMINERALIZATION

Dental caries is a multifactorial disease caused by the interaction of dietary sugars, dental biofilm and the host's dental tissue within the oral environment.¹ It is the cumulative result of consecutive cycles of demineralization and remineralization at

the interface between the biofilm and the tooth surface. Oral bacteria excrete acid after consuming sugar, leading to demineralization.² Upon this acid challenge, the hydroxyapatite crystals are dissolved from the subsurface. Remineralization

Remineralization is the natural repair process for non-cavitated lesions

is the natural repair process for non-cavitated lesions. It relies on calcium and phosphate ions, assisted by fluoride, to rebuild a new surface on the existing crystal remnants in the subsurface. The remineralized crystals are less acid soluble than the original ones.³

Under normal physiological conditions (pH7), saliva is supersaturated with calcium and phos-

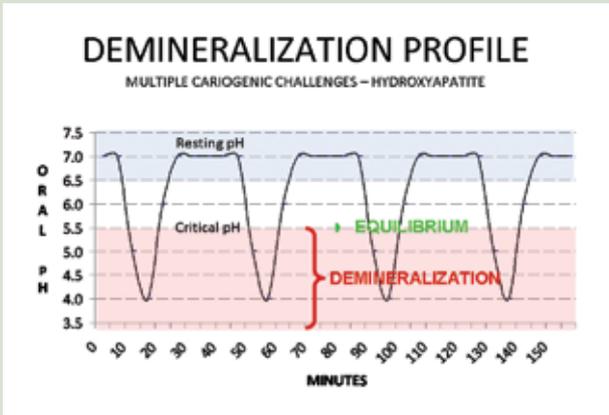


FIGURE 1—Cycling of oral pH during cariogenic challenges in naturally occurring hydroxyapatite.

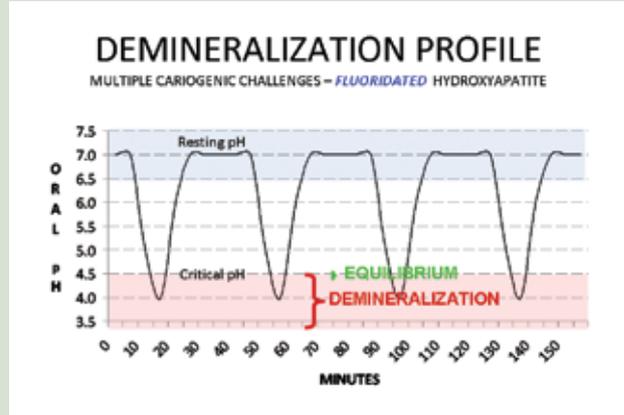


FIGURE 2—Cycling of oral pH during cariogenic challenges in fluoridated hydroxyapatite.

phate ions, making caries progress slow. However, as the bacteria in the biofilm continue to produce acid with sugar consumption, plaque pH falls to 4.5-5.5. This shifts the driving force within the tooth to mineral dissolution.¹ As the pH is lowered, the saturation point of the minerals in the surrounding fluid is changed. The lower the pH, the higher the concentrations of calcium and phosphate required to reach saturation with respect to hydroxyapatite. This is called the “critical pH”, the point where equilibrium exists. There is no mineral dissolution and no mineral precipitation. The critical pH of hydroxyapatite is around 5.5 and that of fluorapatite is around 4.5. This varies with individual patients. Below critical pH, demineralization occurs while above critical pH, remineralization occurs (Figs. 1 & 2).

The critical pH is significantly higher for children than adults. Children have a greater driving force for demineralization in a more acidic oral environment and a decreased driving force for remineralization at normal oral pH. This puts children at greater risk for demineralization than adults.⁴

FLUORIDE

It has been known since the 1980s that fluoride controls caries predominantly through its topical, not systemic, effect.¹ Four mechanisms are involved:

1. Fluoride inhibits demineralization.

If fluoride is present in the plaque fluid when bacteria produce acids, it will penetrate along with the acids at the subsurface, adsorb to the apatite crystal surface and protect the crystals from dissolution.⁵ This coating makes the crystals similar to fluorapa-



tite (critical pH of 4.5) ensuring that no demineralization takes place until the pH reaches this point. Fluoride present in solution at low levels among the enamel crystals can markedly inhibit dissolution of the tooth mineral by acid.^{6,7} This fluoride comes from topical sources such as drinking water, and fluoride products like toothpastes and varnishes. The fluoride, which is incorporated systemically into the tooth, is insufficient to have a measurable effect on its acid solubility.^{7,8}

2. Fluoride enhances remineralization.

When the pH returns to pH 5.5 or above, the saliva which is supersaturated with calcium and phosphate, forces mineral back into the tooth.⁷

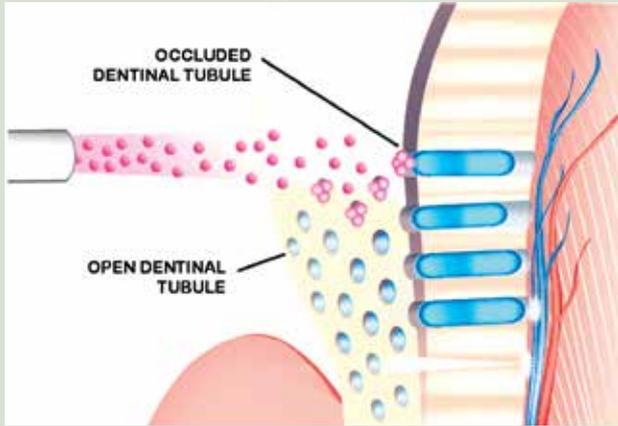


FIGURE 4—NovaMin particles delivered through an air polishing unit clean, desensitize and remineralize to create a smoother less plaque and stain retentive surface.

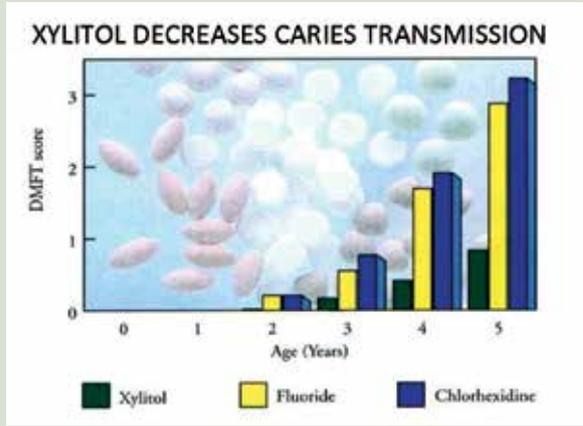


FIGURE 5—Study shows that habitual chewing of xylitol gum by mothers can decrease caries in their children by preventing the transmission of *S. mutans*. Mothers were treated with fluoride varnish, or chlorhexidine rinse or told to chew xylitol gum at 6, 12, and 18 months post-partum.

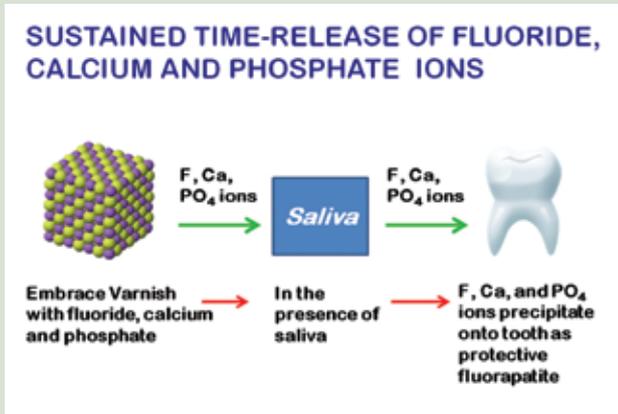


FIGURE 6—cXp technology produces sustained release of calcium, phosphate and fluoride ions over time.

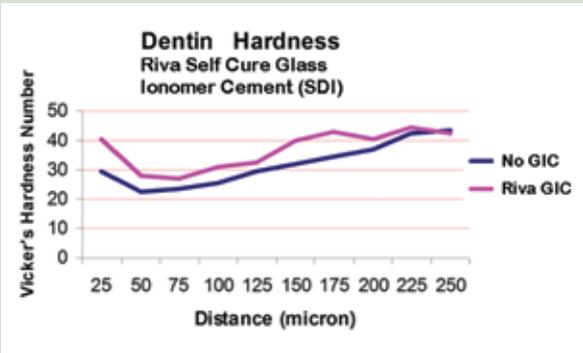


FIGURE 7—Glass Ionomers create an ion enriched harder dentin surface adjacent to the glass ionomer surface.

Children have a greater driving force for demineralization in a more acidic oral environment...than adults

Fluoride adsorbs to the surface of the partially demineralized crystals and attracts calcium ions. This new surface veneer takes up fluoride preferentially from the solution around the crystals and excludes carbonate.⁷

Fluoride speeds up the growth of the new surface by bringing calcium and phosphate ions together

and is also preferentially incorporated into the remineralized surface. This produces a surface which is now more acid resistant.

3. Fluoride may inhibit essential bacterial activity.

Fluoride cannot cross the bacterial cell wall in its ionized form (F⁻). However in an acid environment, F⁻ combines with H⁺ to form HF which easily diffuses into the bacterial cell.^{9,10} Inside the cell HF breaks up and releases fluoride ions that interfere with the essential enzyme activity of the bacterium.

4. Fluoride is retained in intraoral reservoirs after the application of a fluoride treatment such as toothpaste, varnish or restorative material and is then released into the saliva over time.^{11,12}

GIONOMER PRE-REACTED GLASS TECHNOLOGY

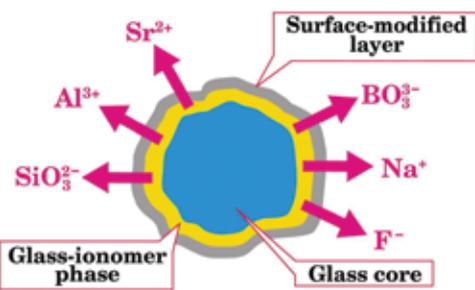


FIGURE 8—Gionomer Pre Reacted Glass (PRG) Technology.

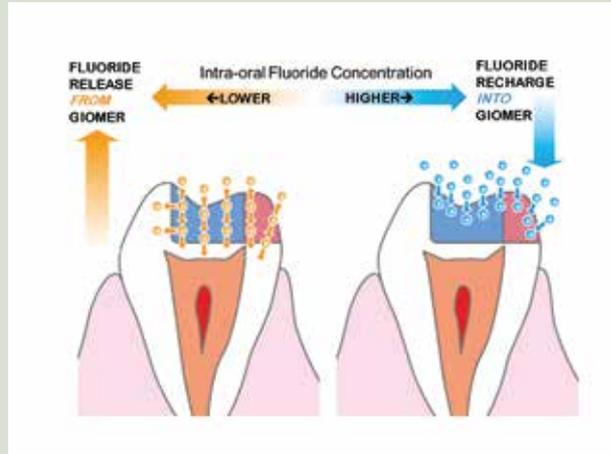


FIGURE 9—Gionomers act as a reservoir to release and recharge fluoride.



FIGURE 10—Biodentine penetrates dentinal tubules forming tag-like structures that create a micromechanical lock with the tooth

The fluoride, which is incorporated systemically into the tooth, is insufficient to have a measurable effect on its acid solubility

Fluoride can remain on dental hard tissue, the oral mucosa or within the dental plaque. Fluoride retention, especially in dental plaque, is clinically beneficial since it can be released during cariogenic challenges to decrease demineralization and enhance remineralization.¹

OTHER REMINERALIZATION THERAPIES

The action of fluoride in remineralization is the gold standard to which newer therapies are compared.

The requirements of an ideal remineralization material are as follows:^{13,14}

- Must diffuse into the subsurface or deliver calcium and phosphate into the subsurface
- Does not deliver an excess of calcium
- Does not favor calculus formation
- Works at an acidic pH
- Works in xerostomic patients
- Boosts the remineralizing properties of saliva
- Shows a benefit over fluoride

The major remineralization technologies that are available in the dental marketplace are Recaldent,

OTHER REMINERALIZATION THERAPIES			
INGREDIENT	DELIVERY	EXAMPLES	
RECALDENT (CPP-ACP)			
	TOPICAL CREAM	GC	MI PASTE
		GC	MI PASTE PLUS
	GUM	TRIDENT	TRIDENT WHITE
NOVAMIN (CALCIUM SODIUM PHOSPHOSILICATE)			
	TOOTH PASTE	ORAL SCIENCE	XPUR TOOTH PASTE
		GSK/ DENTSPLY	SENSODYNE REPAIR & PROTECT
	PROPHY PASTE	DENTSPLY	NUPRO/SENSODYNE PROPHY PASTE
	AIR POLISHING POWDER	ORAL SCIENCE	SYLG POWDER
TCP (TRI-CALCIUM PHOSPHATE)			
	TOOTH PASTE	3MESPE	CLINPRO 5000
	VARNISH	3MESPE	VANISH

TABLE 1

NovaMin, and Tricalcium Phosphate (TCP) (Table 1). They are discussed below:

Recaldent (CPP-ACP)

Recaldent combines phosphoproteins from milk with amorphous calcium phosphate (ACP). ACP on its own simply produces a thin surface coating of hydroxyapatite when applied topically. This is a surface phenomenon that is fundamentally different from the remineralization of enamel subsurface lesions which require the actual penetration of ions into enamel.¹⁴ With the addition of casein phosphopeptide (CPP), Recaldent is more effective than ACP alone.

The CPP in milk stabilizes the calcium and phosphate ions through the formation of complexes which are more readily absorbed by the intestine. The same concept has been applied to Recaldent. The bioavailable complexes of calcium and phosphate are created in the appropriate form for optimal remineralization of subsurface lesions in enamel, not just on the enamel surface. CPP also localizes the ACP in the dental plaque biofilm.¹⁵ The resulting high calcium and phosphate ion concentration gradient drives the ions into the subsurface lesions and achieves high rates of remineralization.¹⁶ Recaldent is available in solutions, gums, lozenges and creams.

NovaMin

NovaMin (GlaxoSmithKline, Brentford, United Kingdom) is technically described as an inor-

ganic amorphous calcium sodium phosphosilicate (CSPS). It belongs to a class of materials which are known as “bioactive glasses”. NovaMin as well as other CSPS materials were originally developed as bone regenerative materials in the early 1970s. Prior to the invention of bioactive glass, all biomaterials were designed to be as inert as possible in the human body.¹⁷ The discovery that a synthetic biomaterial could actually form a chemical bond with bone proved that biomaterials could be engineered to interact with the body. This meant that it was not necessary or advantageous to minimize interactions. It could even be beneficial to encourage them. Bioactive glasses facilitate hydroxyapatite deposition when exposed to fluids containing calcium and phosphate.^{18,19} The mechanism of action is as follows:

In the presence of water or saliva NovaMin rapidly releases sodium ions. This increases the local pH and initiates the release of calcium and phosphate. Many studies have shown NovaMin particles to act as reservoirs and continuously release calcium and phosphate ions into the local environment. This may continue over many days.²⁰ The calcium-phosphate complexes crystallize into hydroxycarbonate apatite, which is chemically and structurally similar to biological apatite.^{21,22} NovaMin has been incorporated into toothpastes, gels and prophy pastes.

A novel delivery system for NovaMin is through an air polishing unit. This procedure has been developed as an improved cleaning method with the added benefit of tooth desensitization and smoothing of surface irregularities

A novel delivery system for NovaMin is through an air polishing unit (Fig. 3). This procedure has been developed as an improved cleaning method with the added benefit of tooth desensitization and smoothing of surface irregularities. Using the Sylic (Oral Science, Montreal, Quebec) technique significantly reduces dentin permeability and completely occlude exposed dentinal tubules.²³ NovaMin powder also has positive remineralization effects on partially and completely demineralized models of dentin. The treatment decreases surface roughness, promoting a smoother, less plaque and stain retentive surface²⁴ (Fig. 4).

AD

Tri-Calcium Phosphate (TCP)

TCP (3M ESPE, London, Ontario) is a bioactive formulation of tri-calcium phosphate and simple organic ingredients. It works synergistically with fluoride to produce superior remineralization of enamel subsurface lesions when compared to using fluoride alone.^{25,26} When it is used in toothpaste formulations, a protective barrier is created around the calcium, allowing it to coexist with the fluoride ions. During toothbrushing, TCP comes into contact with saliva, causing the barrier to dissolve and releasing calcium, phosphate and fluoride. When TCP is incorporated into a 5% NaF varnish, microhardness and acid resistance improve.²⁷ Studies are currently underway to demonstrate the clinical advantages of TCP in rinse form.

...studies have shown that the habitual chewing of xylitol gum by mothers can decrease the caries incidence in their children by preventing the transmission of *S. mutans*

The above remineralization therapies work directly to enhance the concentration of calcium, phosphate and fluoride. The ingredient discussed below, xylitol, works indirectly to promote remineralization by decreasing bacteria and bacterial function and creating the environment where reparative remineralization is optimized.

Xylitol

Xylitol is one of a number of non-sugar sweeteners permitted for use in foods throughout the world. It is found naturally in some foods but is produced primarily from hardwood sources such as birch and beech wood. It is a sugar alcohol that has been shown to have non-cariogenic as well as cariostatic effects.²⁸ More recently it has been shown that the habitual use of xylitol is associated with a significant reduction in caries incidence and increased tooth remineralization.²⁹ Cariogenic bacteria process xylitol very poorly, producing little acid or plaque. This decreases caries incidence and promotes colonization of less virulent strains of bacteria that can ferment xylitol.

Dental caries is an infectious, transmissible, bacterial disease. Most children acquire the bacteria (predominantly *Streptococcus mutans*) from their mothers/caregivers by salivary contact during the emergence of the primary teeth between the ages of 6-30 months.^{30,31} This is called the “discrete win-

dow of infectivity". After the initial colonization of *S. mutans*, the successful establishment of other bacteria on the tooth surfaces is impaired. It has been demonstrated that a reduction of *S. mutans* in the saliva of mothers has resulted in the delayed acquisition of *S. mutans* in their children.^{32,33} And most remarkably, studies have shown that the habitual chewing of xylitol gum by mothers can decrease the caries incidence in their children by preventing the transmission of *S. mutans*³⁴ (Fig. 5).

In fact, chewing xylitol gum decreases caries incidence significantly up to at least five years after the xylitol therapy has been discontinued.³⁵ Children who chew xylitol gum exhibit a significantly lower caries progression and a significant number of caries lesion reversals, suggesting that remineralization has occurred.³⁶ The efficacy of xylitol candies has been shown to be equivalent to that of xylitol gum.³⁷ The dental literature suggests that a minimum of 5-6 grams and three exposures per day (from chewing gum and/or candies) is required for clinical effect.³⁸

A novel method of delivering remineralizing ions (calcium and phosphate) in combination with xylitol has been developed using a NaF varnish (Embrace Varnish, Pulpdent). This varnish contains calcium and phosphate salts that have been nano-coated with xylitol (cXp technology). The xylitol coating prevents early reaction and produces a sustained release of the remineralizing ions. Saliva exposure dissolves the xylitol and frees the calcium and phosphate ions. They then react with the fluoride in the varnish to form protective fluorapatite on the teeth (Fig. 6).

BIOACTIVE RESTORATIVE MATERIALS

When the enamel and dentin no longer have adequate structure to maintain their mineral framework, cavitation takes place and remineralization is an insufficient treatment. Tooth preparation and restoration are now required. Although most restorative materials are inert with respect to the biological tissues of the tooth, some are bioactive. Bioactive restorative materials actually interact with or affect the biological tissues. They effectively work with the dental hard tissues to harden and "heal" them (Table 2). Three bioactive restorative materials are discussed below.

Glass Ionomer Cements

Glass ionomer cements were developed in the early 1970s. They are especially valuable for initial carious lesions, abfractions/erosions/abrasions and for caries control in a high caries risk patients.³⁹

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BIOACTIVE RESTORATIVE MATERIALS

GLASS IONOMER	
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GC	FUJI IX
DENTSPLY	CHEMFIL ROCK
GIOMER	
SHOFU	BEAUTIFIL II HYBRID
	BEAUTIFIL FLOWPLUS FLOWABLE
TRICALCIUM SILICATE CEMENT	
SEPTODONT	BIDENTINE

TABLE 2

Glass ionomers have a true chemical bond with dental tissue. They are bioactive; they encourage remineralization of the surrounding tooth structure and prevent bacterial microleakage through the ion-exchange adhesion that they develop with both enamel and dentin.⁴⁰ This creates a new, ion-enriched material at the tooth-glass ionomer interface. The material consists of phosphate and calcium ions from the dental tissues, and calcium (or strontium), phosphate and aluminum from the glass ionomer cement.⁴⁰ The remineralization process creates a harder dentin surface⁴¹ (Fig. 7). Restoration failure is usually cohesive, leaving the ion exchange layer firmly attached to the cavity wall. The dentinal tubules are sealed and protected from bacterial penetration.⁴²

Fluoride is the catalyst for remineralization, aided by calcium (or strontium) and phosphate. The pattern of release for all the ions in the glass ionomer cement is similar. A low pH actually enhances the process.⁴³

The bioactive remineralizing effect of glass ionomer cements occurs in two distinct regions of the tooth:

1. **The outer surface** of the restoration is exposed to oral fluids and plaque with which it has a continuous exchange of ions.⁴⁴ While wear resistance of the restoration is low at placement, it steadily increases with time and ion uptake.
2. **The inner surface** of the restoration, adjacent to the preparation, is isolated from the oral environ-

ment. The continuous flow of dentinal fluid creates a wet environment that is conducive to the exchange of ions. At placement, there is a significant release of ions from the cement which combine with similar ions from the dentinal fluid to promote remineralization. After the glass ionomer sets, there is a continuous low-level ion exchange which accounts for the remineralization of the tooth surface that is found clinically.⁴²

Giomers

Giomers (Shofu Dental, San Marcos, CA) are the latest category of hybrid restorative materials and they are bioactive as well. Giomer technology represents the true hybridization of glass ionomers and composite resins. There is an ideal combination of the properties of these two distinct restorative categories: the fluoride release and recharge of glass ionomers and the esthetics, physical properties and handling of composite resins.⁴⁵

The Giomer concept is based on PRG (Pre Reacted Glass) technology: a glass core, surrounded by a glass ionomer phase enclosed within a polyacid matrix (Fig. 8). Studies show dentin remineralization occurs at the preparation surface adjacent to the giomer.⁴⁶

The ions within the pre reacted glass particles have distinct biological effects. The fluoride, as discussed above, improves acid resistance through the formation of fluorapatite, remineralizes decalcified tooth substance and is antibacterial. The strontium ion improves acid resistance by forming stontiumapatite, inhibits dentin hypersensitivity and has been shown to accelerate the formation of bone. The aluminum ion inhibits dentin hypersensitivity. The silicate ion aids in the calcification of bone while the borate ion accelerates bone formation and is bactericidal.

Giomer technology represents the true hybridization of glass ionomers and composite resins

Giomers are also able to take up extra fluoride (after fluoride toothpaste, rinse or varnish are applied) from the oral fluids and then act as a reservoir until the fluoride is needed. This is called “fluoride release and recharge” (Fig. 9). Giomers release and recharge fluoride efficiently and significantly better

than compomers⁴⁷ and composite resins although not as well as glass ionomers.⁴⁸ Gioner fissure sealants have superior recharge and release of ions when they are compared to resin sealants. Hence, they work actively to decrease demineralization and increase remineralization in young teeth at their most caries susceptible stage.⁴⁹

Gioners resist plaque formation.⁵⁰ A “material film layer” forms on the surface of the gioner with salivary contact. It consists of aluminum, silica, strontium and other ions which originate from the PRG fillers and act to inhibit bacterial adhesion. The clinical performance of gioners has been tested against those of high quality hybrid resin composites. Gioners have been found to compare favorably for all criteria.⁵¹

Biodentine has the potential to heal pulps, avoiding what may have been inevitable endodontic treatment in the past

Biodentine Tricalcium Silicate Cement

Biodentine (Septodont, Cambridge, Ontario) is a new bioactive calcium silicate based product that has been designed as an all-around “dentin replacement” material. It can be used in endodontic repair (root perforations, apexification, resorptive lesions), pulp capping, as well as a dentin replacement in restorative dentistry. It was formulated by taking the MTA-based endodontic repair cement technology, improving its physical and handling properties, and creating a dentin replacement material with significant reparative qualities.

Biodentine penetrates the dentinal tubules forming tag-like structures that create a micromechanical lock with the tooth. It then begins to stimulate reparative dentin (Fig. 10).

Biodentine has been shown to enhance the formation of reparatory dentin and to create a dense dentin barrier after direct pulp capping^{52,53} as well as healing damaged pulp fibroblasts.⁵⁴ Clinical trials confirm Biodentine’s ability to preserve pulp vitality even in very difficult cases. Biodentine has the potential to heal pulps, avoiding what may have been inevitable endodontic treatment in the past.

CONCLUSION

The medical model of Proactive Intervention is be-

coming the paradigm in dental care. This should be an integral part of daily practice and not relegated to the “preventive” side of the office. The multifactorial disease process of demineralization and caries can be slowed or even stopped before more extensive treatment becomes necessary. The oral care provider has simple remineralization tools, techniques and products that have been found effective in reversing and controlling the caries process. They can and should be used proactively to maintain hard tissue health throughout the patient’s life.

Dr. Fay Goldstep has served on the teaching faculties of the Post-graduate Programs in Esthetic Dentistry at SUNY Buffalo, Baylor, Universities of Florida (Gainesville), Minnesota (Minneapolis), California (UCSF) and UMKC (Kansas City). She has lectured on Proactive Intervention Dentistry and Lasers as Soft Tissue Handpieces nationally and internationally, including the ADA, PDC, CDA, Yankee, AACD, AGD, and the Big Apple dental conferences. Dr. Goldstep is on the Editorial Board of Oral Health Magazine (Healing/Preventive Dentistry) and Dental Tribune US Edition. She is a Fellow of the American College of Dentists, International Academy of Dental-Facial Esthetics and the Academy of Dentistry International. Dr. Goldstep has been a contributing author to 4 textbooks and has published more than 25 articles. She has been listed as one of the leaders in Continuing Education by Dentistry Today since 2002. Dr. Goldstep is a consultant to a number of dental companies and maintains a private practice in Markham, Ontario.

Oral Health welcomes this original article.

REFERENCES

1. Buzalaf MAR, Fluoride and the Oral Environment, Monogr Oral Sci, Basel, Karger, 2011, vol 22, p 97-114.
2. Fejerskov O, Kidd, EA, Nyvad B, Baclum V, Defining the disease: an introduction: in Fejerskov O, KiddE (eds): Dental Caries- The Disease and its Clinical Magement, ed 2, Oxford, Blackwell Munksgaard, 2008, p 3-6.
3. Featherstone JD, Dental Caries: a dynamic disease process, Aust Dent J, 2008;53(3):286-91.
4. Anderson P, Hector MP, Rampasad MA, Critical pH in resting and simulated whole saliva in groups of children and adults, Int J Paediatr Dent, 2001, 11 (4):266-73.
5. Featherstone JDB, Prevention and reversal of dental caries: role of low level fluoride, community Dent Oral Epidemiol 1999, 27: 31-40.
6. Featherstone JDB, Glena r, Shariati M, Shields CP, Dependence of invitro demineralization and remineralization of dental enamel on fluoride concentration, J Dent Res, 1990, 69:620-5.
7. Ten Cate JM, feathersone JDB, Mechaanistic aspects of the interactions between fluoride and dental enamel, Crit Rev Oral Biol 1991, 2:283-96.
8. Fejerskov O, Thystrup A, Larsen MJ, Rational use of fluorides in caries prevention, ACTA Odontol Scand, 1981:39:241-9.
9. Hamilton IR, Bowden GHW, Fluoride effects on oral bacteria, In Fejerskov O Ekstrand J, Burt BA (eds) Fluoride in Dentistry, Copenhagen, Munksgaard, 1996 p 230-51.
10. Van Louveren C, the antimicrobial action of fluoride and its role in caries inhibition, J Dent Res1990:69:676-81.
11. Whitford GM, Wadison JL, Schafer TE, Adair SM, Plaque fluoride concentrations are dependent on plaque calcium concentrations, Caries Res 2002, 36:256-265.
12. Pessan JP, Alves KM, Ramires I, et al, Effects of regular and low-fluoride, J Dent Res, 2010, 89:1106-1110.
13. Zero DT, Dentifrices, mouthwashes and remineralization/caries arrestment strategies, BMC Oral Health, 2006:6 (Suppl 1):S9-S22.
14. Walsh L, Evidence that demands a verdict: latest developments in remineralization

- therapies, Dental Economics, 2009.
15. Cross KJ, Huq NL, Reynolds EC, Casein Phosphopeptides in oral health-chemistry and clinical applications, Curr Pharm Des, 2007, 13 (8):793-800.
 16. Reynolds EC, Remineralization of enamel subsurface lesions by casein phosphopeptide-stabilization calcium phosphate solutions, J Dent Res, 1997;79(9):1587-95.
 17. Hench L Biomaterials, Science, 1980:208:826-831
 18. Burwell AK, Litkowski LJ, Greenspan DC, Calcium sodium phosphosilicate (Novamin(r)): remineralization potential, Advances in Dental Research, 2009, 21:35-9
 19. Anderson OH, Kangasniemi I, Calcium phosphate formation at the surface of bioactive glass in vitro, Journal of Biomedical Material Research, 1991, 25:1019-30.
 20. Damen JJ, ten Cate JM, Silica-induced precipitation of calcium phosphate in the presence of inhibitors of hydroxyapatite formation, J Dent Res, 1992: 71:453-457.
 21. Gandolfi MG, Silvia F, H PD, Gasparotto G, Carolo P, Calcium silicate coating derived from Portland cement as a treatment for hypersensitive dentine, Journal of Dentistry, 2008,36(8),565-578.
 22. Anderson OH, Kangasniemi I, Calcium phosphate formation at the surface of bioactive glass in vitro, Journal of Biomedical Materials Research, 1991, 25(8), 1019-1030.
 23. Sauro, S, Watson T, Thompson I, Ultramorphology and dentine permeability changes induced by phophylactic procedures on exposed dentinal tubules in middle dentine, Med Oral Patol Oral Cir Buccal, Biomaterials and bioengineering in dentistry, 2011.
 24. Wang Z, et al, Dentine remineralization induced by two bioactive glasses developed for air abrasion purposes, Journal of Dentistry, 2011, doi:10.1016/j.dent.2011.08.006.
 25. Karlinsky RL, Mackey AC, Walker ER, Frederick KE, Enhancing Remineralization of Subsurface Enamel Lesions with Functionalized FTCP, In Biomaterials Developments and Applications, 2010, EDS H Bourg, A Lisle:353-374.
 26. Karlinsky RL, Mackey AC, Solid-State Preparation and Dental Application of an Organically Modified Calcium Phosphate, J Mater Sci, 2009:44:346-349
 27. Flanigan PJ, Vang F and Pfarrer AM, Remineralization and Acid Resistance Effects of 5% NaF Varnishes, J dent Res 89 (Spec Iss B), 383, 2010.
 28. Maguire A, Rugg-Gunn AJ, Xylitol and caries prevention—is it a magic bullet?, British Dental Journal, 2003:194:429-436.
 29. Makinen, K, Sugar Alcohols, Caries Incidence and Remineralization of Caries Lesions : A Literature Review, International Journal of Dentistry, Vol 2010, Article ID 981072, 23 pages, doi 10.115/2010/981072.
 30. Berkowitz, RJ, Turner J, Green P, Maternal salivary levels of Streptococcus mutans and primary oral infection of infants, Arch Oral Biol, 1981, 26:147-149.
 31. Caufield PQ, Cutter GR, Dasanayake AP, Initial acquisition of mutans streptococci by infants: evidence for a discrete window of infectivity, 1993, J Dent Res, 72:37-45.
 32. Kohler B, Andreen I, Influence of caries-preventive measures in mothers on cariogenic bacteria and caries experience in their children, 1994, Arch Oral Biol, 39:907-911.
 33. Kohler B, Bratthall D, Krasse B, Preventive measures in mothers influence the establishment of the bacterium, Streptococcus mutans in their infants, Arch Oral Biol, 28:225-231.
 34. Isokangas P, Soderling E, Pienihakkinen K, Alanen P, Occurrence of Dental Decay in Children after Maternal Consumption of Xylitol Chewing gum, a Follow up from 0 to 5 Years of Age, Journal of dental Research, 2000,79,1885.
 35. Isokangas P, Makinen KK, Tiekso J, Alanen P, Long term effect of xylitol chewing gum in the prevention of dental caries: a follow up 5 years after termination of a prevention program, Caries Res, 1993:27:495-498.
 36. Kandelman D, Gagnon G, Clinical results after 12 months from a study of the incidence and progression of dental caries in relation to consumption of chewing gum containing xylitol in school preventive programs, J Dent Res, 1987:66:1407-1411.
 37. Alanen P, Isokangas P, Gutmann K, Xylitol candies in caries prevention: results of a field study in Estonian children, Community Dent Oral Epidemiol, 2000: 28:218-224.
 38. Milgrom P, Ly KA, Rothen M, Xylitol and Its Vehicles for Public Health needs, Advances in Dental Research, 2009, doi:10.1177/089593740093335623.
 39. Mount GJ, An Atlas of Glass-Ionomer Cement: A Clinicians Guide, 2nd Ed, London, Martin Dunitz, 1994.
 40. Mount GJ, Adhesion of glass-ionomer cement in the clinical environment, Oper Dent, 1991, 16:141-148
 41. McIntyre J.M., Cheetham J., Dalidjan M.; Ionic Exchange Between Riva Self Cure GIC and Demineralized Dentine; Brisbane 2006 IADR Abstract #2078, University of Adelaide, Australia.
 42. Mount, G, Advances in Glass Ionomer Cements—Chapter 14, Glass Ionomers: Advantages, Disadvantages, and Future Implications, Quintessence Publishing, 1999, p269-293.
 43. Ngo H, Marino V, Mount GJ, Calcium strontium, aluminum, sodium and fluoride release from four glass-ionomers, J Dent Res, 1998:77:641 (abstract 75).
 44. De Moor RJG, Verbeek RM, De Meeyer EAP, Fluoride release profiles of restorative glass ionomer formulations, Dent Mater 1996:12:88-95.
 45. Koirala S, Yap A, A Clinical Guide to Direct Cosmetic Restorations with Giomer, Dental Tribune International, 2008.
 46. Miyauchi T et al, The effect of Giomer restorative materials on demineralized dentin, 2010, IADR Abstract 135006.
 47. Dhull KS, Nandlal B, Comparative evaluation of fluoride release from PRG-composites and compomer on application of topical fluoride, Journal of Indian Society of Pedodontics and Preventive Dentistry, 2009:27:1:27-32.
 48. Okuyama K, et al, Fluoride release and uptake by various dental materials after fluoride application, Am J Dent, 2006:19:123-127.
 49. Shimazu Kasaki et al, Evaluation of the ion-releasing and recharging abilities of a resin-based fissure sealant containing S-PRG filler, Dental Materials Journal, 2011:30(6):923-927.
 50. Honda T, Yamamoto K et al, Study on the firm substance produced from S-PRG filler, JJ Conser Dent, 2002:45(Autumn):42.
 51. Tian FC, Clinical performance of Giomer restorative system, 2010 IADR Abstract.
 52. About I, Coiffage pulpaire direct de RD94 a l'aide du modele de culture de dent entire, Report RD EN RA EXT-RD94/096, 2007.
 53. Shayegan A, RD 94 Etude #PC08-001, Etude de RD 94 comme agent pulpaire dans le cadre de pulpotomie et coiffage direct sur les dents lacteales de cochon, Report RD RA DEV 94-006, 2009.
 54. About I, Effects des materiaux bioactifs Biodentine (tm) et Calcipulpe(r) sure les etapes precoces de la regeneration dentaire, Report RD RA DEV 94-013, 2009.

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