

Secondary Caries and Microleakage

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Alligator #1, #2 & #3
Gainesville, Florida, 2009

Secondary caries –questions


1. How to best predict secondary caries?
2. How to best prevent secondary caries?
3. How to best identify secondary caries (early)?
4. How to best manage secondary caries?

Secondary caries –questions

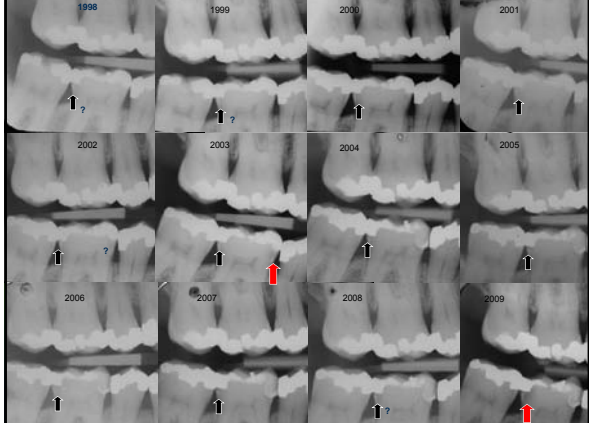
The answers will undoubtedly be influenced by the **stakeholders' understanding of etiopathogenesis**

1. How to best predict secondary caries?
2. How to best prevent secondary caries?
3. How to best identify secondary caries (early)?
4. How to best manage secondary caries?

Etiopathogenesis of secondary caries – 1/3



Bulk fracture → caries OR Caries → bulk fracture?



Etiopathogenesis of secondary caries – 3/3

130 International Dental Journal (1992) Vol. 42/No. 3

Figure 5
 • A diagrammatic representation of secondary caries showing that the lesion may occur in two parts: an 'outer lesion' formed on the surface of the tooth as a result of primary attack and a 'cavity wall lesion' formed as a result of leakage between the restoration and the cavity wall.

Kiidi, Dental Update 1981
 Kidd, Quintessence Publ., Co. 1989
 Kidd, Toffenetti & Mjör. Int Dent J 1992

Secondary caries –questions

The answers will undoubtedly be influenced by the **stakeholders' understanding of etiopathogenesis**

- How to best predict secondary caries?
- How to best prevent secondary caries?
 Material – operator – patient factors
- How to best identify secondary caries (early)?
 - Understanding of etiopathogenesis + Diagnostic test validity and reliability
- How to best manage secondary caries?
 - Understanding of etiopathogenesis + Effectiveness of interventions

Desire to avoid adverse clinical outcomes by the development of (minimum) specifications for dental materials

1900 1920 1930 1940 1950 1960

ADA fdi

Desire to avoid adverse clinical outcomes by the development of (minimum) specifications for dental materials

1920 1930 1940 1950 1960 1970 1980 1990 2000

ADA - ASTM / ANSI ... BSI ... DIN ... AFNOR ... NIOM ... Australia DMRL ...

A listing of the current specifications for dental materials throughout the world shows the following:

Organization	Number of Specifications
Federation Dentaire Internationale (FDI)	Uses ISO Standards
International Organization for Standardization (ISO)	24
American Dental Association (ADA)	65
Dental Specifications of Various States of the U.S.A.	10
American National Standards Institute (ANSI)	34
Defense Medical Material Board (DMMB) (U.S.A.)	33*
Standards Association of Australia (SAA)	54
British Standards Institution (BSI)	30
Canadian Standards Association (CSA)	9
Czechoslovakian Society for Standardization and Measuring (CSN)	7
French Standards Association (AFNOR)	29
German Standards Institute (DIN)	60
Hungarian Office of Standardization (MSZH)	27
Indian Standards Institution (ISI)	155
Standards Institution of Israel (SII)	3
Japanese Standards Association (JIS)	63
Polish Committee of Standardization and Measures (PKM)	45
South African Bureau of Standards (SABS)	21
Swedish Planning and Rationalization Institute (SRI)	63

From: Paffenbarger, et al. US. National Bureau of Standards pub. #571, 1980

Specifications according to tests – which ones predict adverse clinical outcomes?

Static tests ?
 Compressive (crushing) strength, e.g., 1h. & 24 h.
 Tensile strength, e.g., 5 min.
 Transverse strength, e.g., 1h. & 24 h.
 (Flexure/bending/modulus of rupture)
 Modulus of elasticity (Young's Modulus)
 Shear modulus

Dynamic tests ?
 Compressive modulus
 Tensile modulus
 Bending modulus
 Resilience
 Fatigue
 Fracture toughness

Validity of a test

- Reproducible
- Known parameters
- Low C.V. (#samples)
- Calibrated devices

Specifications according to tests – which ones predict adverse clinical outcomes?

Other defined tests?
 Flow (Creep), 3-24 h.
 Dimensional change, e.g., 5 min. -24 h.
 Polymerization- /Setting-...contraction/expansion
 Hardness
 Thermal expansion coefficient
 Water solubility / - sorption

Other undefined tests?
 Abrasion resistance (Wear)
 Adhesion
 Surface roughness
 Marginal leakage
 «Retention strength»
 Color stability

Validity of a test

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«Neither dentists nor laboratory researchers have a clue as to what these tests say on possible clinical outcome in terms of predictability and longevity»

Dr. Siegwad D. Heintze, Head of Preclinical Research, Ivoclar Vivadent. Dent Mater 2013.

Evidence that specifications according to tests predict adverse clinical outcomes?

Weak - according to current leading content experts, although perhaps substantially inferior products may be identified

Articles shown include:
 - In Vitro Performance of Class I and II Composite Restorations: A Literature Review on Nondestructive Laboratory Trials—Part I
 - Clinical relevance of tests on bond strength, microleakage and marginal adaptation
 - Correlation of clinical performance with 'in vitro tests' of restorative dental materials that use polymer-based matrices
 - Review Article: Prediction and diagnosis of clinical outcomes affecting restoration margins
 - Systematic Reviews: I. The Correlation Between Laboratory Tests on Marginal Quality and Bond Strength; II. The Correlation Between Marginal Quality and Clinical Outcome
 - Resin-based composite performance: Are there some things we can't predict?

Restoration material and adverse outcome undesirable performance

- Degradation
 - Bulk (/surface)
 - Interface (/margin)
- Material loss
- Fractures/cracks
- Rough surface
- Poor adaptation to tooth tissues
- Discoloration
 - Surface (/bulk)
 - Margin (/interface)

What are the predictors?

Restoration material and adverse outcome undesirable performance versus risk factor for:

- Degradation
 - Bulk (/surface)
 - Interface (/margin)
- Material loss
- Fractures/cracks
- Rough surface
- Poor adaptation to tooth tissues
- Discoloration
 - Surface (/bulk)
 - Margin (/interface)

- Secondary caries
- Fractures
- Hyper-sensitivity
- Pulpal injury
- Antagonist extrusion
- Impaction/periodontal disease
- Shade

What are the predictors? Why and how much are the risks inflated?

Desire to avoid adverse clinical outcomes by the development of standards for clinical practice and research

1920 1930 1940 1950 1960

ADA

- 1971: Cvar & Ryge, "Ryge system" US Dept. Health, Educ. & Welfare
- 1972: ADA Recommended standard practices for clinical evaluation of dental materials and devices
- 1973: ADA Guidelines for reporting clinical trials
- 1977: California Dental Association "CDA system"
- 1978: Clinical evaluation of dental materials. US Dept. Health & H.S.; 1980 - "USPHS system"

Desire to avoid adverse clinical outcomes by the development of standards for clinical practice and research

1920 1930 1940 1950 1960 1970

ADA

fdi

- 1977: Recommended format. Clinical comparison of several anterior and posterior materials.
- 1982: Recommendations for clinical research protocols for dental materials
- 1990: Good manufacturing practices, including quality assurance for dental materials
- 1981: Expansion of the ADA acceptance program. Composite resin materials for occlusal class I and II restorations
- 1986: Evaluation of dentin and enamel adhesive materials (r1991, r1994, r2001)
- 1989: Composite resins for posterior restorations (r1996, r2001)

Desire to avoid adverse clinical outcomes by the development of standards for clinical practice and research

2007: Hickel et al. Recommendations for conducting controlled clinical studies of dental restorative materials & criteria for evaluation of direct and indirect restorations including onlays and partial crowns.
2010: Hickel et al. Clinical criteria for the evaluation of direct and indirect restorations. Update

ISO/TC194 Clinical investigation of medical devices for human subjects - Good clinical practice

1920 1930 1940 1950 1960 1970 1980 1990 2000

ISO TC 194 Dentistry

SC1 Filling and restorative materials: 14 wgs
SC2 Prosthodontic materials: 20 wgs
SC3 Terminology: 4 wgs
SC4 Dental instruments: 10 wgs
SC6 Dental equipment: 8 wgs
SC7 Oral hygiene products: 4 wgs
SC8 Dental implants: 5 wgs
SC9 CAD/CAM: 4 wgs

Only test standards

ADA 2001-2006, Acceptance program guidelines for resin-based composites for posterior restorations & for dentin and enamel adhesive materials

Topics discussed in this presentation

- Dental caries, a brief review
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DENTAL CARIES, A BRIEF REVIEW

WHAT IS DENTAL CARIES?

Manifested in dental hard tissues by:

1. demineralization
2. white OR brown (spot) lesion
3. cavity ("cavitation")

Specific/Colloquial terms

- active caries
- arrested caries
- caries lesion
- cavity
- chemical dissolution by microbes
- decay
- demineralization
- histologic caries
- inactive caries
- infected dentin
- radiographic caries

Enamel demineralization (Microscopically*)

- 1 Surface zone
- 2 Lesion body zone
- 3 Dark zone
- 4 Translucent zone

(*change of optical birefringence)

Dentin sclerotization

Microbial biofilm i.e., «plaque»

time

«Cavitation»

Lesion width in dentin ≤ enamel* (*some exceptions)

Central transversal line

Cellular changes in the pulp

Tertiary dentin in the pulp

Microradiograph: de Medeiros et al. J Microscopy 2012

Histopathology of enamel caries

Changes within the zones of a caries lesion

Relative pore structure of the 4 caries lesion zones

Relative concentrations of important ions. Selective loss of Mg and carbonate is illustrated together with concentration gradients of fluoride and protons from surface to interior.

Proposed phase changes in the surface zone and the positively birefringent zone following ingress of protons and fluoride and a net loss of mineral.

Net chemical changes detected at each stage of carious attack.

Diagrammatic representation of changes in enamel mineral crystal morphology within each zone to account for changes in pore structure.

Source: Robinson et al. 2000

Mineral loss detection methods

Laboratory methods

Destructive methods

- Chemical analysis
- X-sectional microhardness
- Optical birefringence (Polarized light)
- Confocal light microscopy
- Laser scanning microscopy (CLSM)
- Transverse microradiography (TMR)
- Microprobe analysis
- Energy-dispersive spectroscopy (SEM-EDX)
- Raman spectroscopy, and Fourier-transform infrared spectroscopy (FTIR)

Non-destructive sequential methods

- Surface microhardness
- Iodine absorptiometry / penetration
- Longitudinal MR (LMR)
- Light scattering
- Wavelength-independent MR (WIM_{IR-WIM})

Intraoral methods

- Light-, infrared- or laser-induced fluorescence
- Electrical conductivity
- Computerized radiography +/- algorithms for automated detection of lesions
- Optical coherence tomography (OCT)
- Polarization-sensitive OCT (PS-OCT) in combination with near-infrared light

Variability

Discrimination threshold in dental tissues, resolution, time, costs & complexity

Enamel, Optical Birefringence* & Imbibition Media**

* Intrinsic (AKA crystalline) o.b. or anisotropy to light & Form (AKA textural or structure) o.b. or anisotropy to light
 ** Effects depend on refractive index & molecular weight

	Quinoline transmitted light	Thoulet sol. transmitted light		Micro-radiograph	Water
	Quinoline polarized light 45°	Thoulet sol. polarized light 45°		Air	Dehydrate w/ ethyl alcohol & air
	Canada balsam transmitted light	Canada balsam polarized light 45°		Quinoline	
	Quinoline transmitted light	Quinoline polarized light 45°			
	Quinoline transmitted light	Canada balsam transmitted light			

Source: Darling et al. Arch Oral Biol 1961
 Source: de Medeiros et al. J Microscopy 2012

Creating caries-like lesions artificially in vitro

- Human and non-human, mainly bovine, teeth or tooth specimens
- Several approaches, some specifically tailored to create lesions in enamel or in dentin or in root cement.
- Two methods prevail, (1) an acidified medium with/without buffering & with/without pH-cycling; (2) an acidified broth containing usually some strain of *Streptococcus mutans*
- Mineral loss profiles of the surface and subsurface zones differ with method
- Differences can be large, e.g., at pH=5, a carboxymethyl cellulose gel (6%) causes ~33 volume % mineral loss in enamel per day, while an unstirred solution causes 13% & 26% volume % mineral loss in enamel per hour (with & without added fluorides respectively).
- The ultimate hope is to build the artificial mouth, or at least a steady state microcosm. In spite of some elaborate contraptions we have not succeeded yet to simulate the complexities of the intraoral ecology and microenvironment

Creating caries-like lesions artificially in situ

- Since early 90'ies. Used for multiple research objectives, e.g., assessing erosive or cariogenic potential of various substances, or, appraising the potential for remineralization following application of various oral care products on preconditioned specimens
- Human and non-human specimens
- Specimens mounted in a dental device worn by subjects for various periods
- Surface of the specimens often covered or machined to increase plaque retention
- Deminerzalization/erosion accelerated by repeat bathing of the device, e.g., caries, 4-8x / day in a 20% sucrose solution
- Deminerzalization differs from in vitro setups

Source: Arends et al. Adv Dent Res 1997

Are there any differences?

Artificial *caries-like* lesion
 versus
 Artificial *caries* lesion
 versus
 Clinical caries lesion

How is the disease entitled «Dental caries» defined by laypersons, basic scientists, dentists, clinical researchers and epidemiologists?

What is dental caries?

Essentialism and nominalism in medicine: logic of diagnosis in disease terminology

J.G. Scadding

Understanding of disease concept:
Essentialistic: The disease in itself exist, but our criteria for describing the disease change over time
Nominalistic: The actual signs and symptoms constitute the disease

FOCUS
 Dental caries paradigms in diagnosis and diagnostic research

Åslund V, Heimdann J, Nyvad B. Dental caries paradigms in diagnosis and diagnostic research. Eur J Oral Sci 2006; 124: 203-211. © 2006 The Authors. Journal compilation © 2006 Eur J Oral Sci

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Restoration:tooth interface morphologies on non-occluding tooth surfaces

Interface descriptors: «Margin(-al)» - «adaptation» - «gap». Other terms are more specifically linked to:
Adhesive materials: «seal(-ing)», Amalgam:«crevice»/«ditch», Indirect restorations: «discrepancy»/«fit»

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Is it possible that the choice of term to describe the restoration-tooth interface influence clinicians' treatment decisions (perhaps unwittingly)?

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Interfaces are never static, but in a complex dynamic equilibrium with substances and minerals in the saliva and in the more or less porous hard tissues of the tooth

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Interfaces are never static, but in a complex dynamic equilibrium with substances and minerals in the saliva and in the more or less porous hard tissues of the tooth

Can fatigue degradation of the interface be simulated - influenced by compressive, tensile and shear stresses, compressive, tensile and amounts, salivary composition and amounts, shifting occlusal loads, dietary habits and oral cleanliness and possibly the individual's intra-oral bacterial profile - and will one test likely fit all materials?

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Etiopathogenesis of secondary caries – which one prevails?

The origin of the term «cavity wall lesion*»

*Hals & Nernæs, Caries Res 1971

Amalgam (in vivo/vitro 1975)

Resin composite (in vitro, 1976)

M&M: Polarized light microscopy x60 & Quinoline imbibition

Cavity wall lesion: "tending to encompass filling..." "usually without penetrating deeply into the tissue"

The depiction of the «wall lesion» originally

Fig. 12. Tentative explanation of radiopaque layer of enamel cavity wall. Large arrow indicates progression of decalcification process. Small arrows indicate attack of hydrogen ions (H⁺) on enamel surface and enamel cavity margin. Reaction products of decalcification process (R, backward arrows) will precipitate in enamel surface and may do so in the enamel cavity margin as well.

From: *Hals & Nernæs, Caries Res 1971

A tentative explanation for a radiopaque layer of the cavity wall

The original and the modified «wall lesion»

Depictions of a radiopaque layer of the cavity wall in original publications

A modified redrawing appearing ~10 years later - allegedly depicting "secondary caries"

From: Hals & Nernæs 1971, Hals & Halse Hals & Laegreid 1975, 1976

Figure 3: A diagrammatic representation of secondary caries showing that the lesion may occur in two parts: an "outer lesion" formed on the surface of the tooth as a result of primary attack and a cavity wall lesion formed as a result of leakage between the restoration and the cavity wall.

Kidd, Dental Update 1981
Kidd, Quintessence Publ., Co. 1989
Kidd, Toffenetti & Mjør, Int Dent J 1992

The end of the faith in «wall lesions»

An etiopathogenetic theory which persisted for a decade before a "correction" was made in a 2003 cariology textbook

2003, 2008, 2015

(pp. 272): "...gaps will facilitate pathways for microorganisms, but this does not mean that these cause a caries reaction deep within the gaps, as was once believed. It should also be kept in mind that these are not empty spaces. They will be filled with proteinaceous material from dentin liquid and saliva."

The re-emergence of use of the term «wall lesion» in laboratory experiments

The extent of demineralization as an effect of the distance between a specimen and a block of material placed in an acidified medium

Derand et al. Swed Dent J 1991

Totiam et al. Caries Res 2007

Diercke et al. Clin Oral Invest 2009

Nassar & Gonzalez-Cabezas Caries Res 2011

The re-emergence of use of the term «wall lesion» in in situ experiments

Lennon et al. Eur J Oral Sci 2007

Thomas et al. Caries Res 2007 & Oral Microbiol Immunol 2008

Van de Sande et al. J Dent 2014

Kuper et al. J Dent Res 2014 & Montagner et al. J Dent 2015

1.6 x 3 x 1.5mm

1.5x2x1mm

Specimen size

What is dental caries?

Essentialism and nominalism in medicine: logic of diagnosis in disease terminology

J.G. Scadding

Medical discourse is essentially conducted with a terminology which implies that diseases are causes of illness. The pathology of this discourse has long been evident – and because obvious when always some made an explicit reference to diagnosis. Controversy about definitions of disease restores.

Understanding of disease concept:
Essentialistic: The disease in itself exist, but our criteria for describing the disease change over time
Nominalistic: The actual signs and symptoms constitute the disease

FOCUS
Dental caries paradigms in diagnosis and diagnostic research

Vibeke Børlum, Jona Heidmann*, Bente Nyvad**

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Ashwin T. Meuwagen, J. Snydal B. Dental caries paradigms in diagnosis and diagnostic research. Eur J Oral Sci 2008; 114: 263-277. © 2008 The Authors. Journal compilation © 2008 Eur J Oral Sci

In this article, the fundamentals of caries diagnosis are reviewed from three component perspectives, namely the strategy, the logic, and the tactics. Strategy concerns the objectives of the diagnostic process (i.e. why we diagnose caries). The logic describes how we assemble and evaluate the information collected and how this leads to an assessment of diagnostic value. Finally, tactics are about how we collect the information relevant to arrive at a correct diagnosis. We argue that the **holistic, essentialistic paradigm of disease should be replaced by a nominalistic paradigm**. This allows us to overcome the problem of a lack of a correct gold standard and to proceed to caries-diagnostic research that the diagnostic methods that result in the best health outcomes for our patients. We also demonstrate the limitations of the medical model when attempting to understand caries diagnosis, and adhere to the

The risk of misunderstanding when a poor choice of the term «wall lesion» is combined with an essentialistic perspective on caries

Essentialistic: The disease in itself exist, but our criteria for describing the disease change over time

Dis Oral Investis. 2009 Dec; 13(4):439-44. doi: 10.1007/s00714-009-0210-z. Epub 2009 Feb 12.

Isolated development of inner (wall) caries like lesions in a bacterial-based in vitro model.

Diercke S, Lussi A, Skarsten T, Seemann B*

Author information

Abstract
 The study conducted in a bacterial-based in vitro caries model aimed to determine whether typical inner (secondary) caries lesions can be detected as cavity walls of restorations with selected gap widths when the development of outer lesions is minimized. Thirty bovine tooth specimens were randomly assigned to the following groups: test group 50 (TG50; gap, 50 microm), test group 100 (TG100; gap, 100 microm), test group 250 (TG250; gap, 250 microm) and a control group (CG; gap, 250 microm). The outer tooth surface of the test group specimens was covered with an acid-resistant varnish to inhibit the development of an outer caries lesion. After incubation in the caries model, the **presence of demarcation lines** at the cavity wall was determined by confocal laser scanning microscopy. All test group specimens demonstrated only wall lesions. The CG specimens developed outer and wall lesions. The TG250 specimens showed significantly less wall lesion area compared to the CG ($p < 0.05$). In the test groups, a statistically significant increase ($p < 0.05$) in lesion area could be detected in enamel between TG50 and TG250 and in dentine between TG50 and TG100. In conclusion, the inner-wall lesions of secondary caries can develop without the presence of outer lesions and therefore can be regarded as an entity on their own. The extent of independently developed wall lesions increased with gap width in the present setting.

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Overview of some experimental variables

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Further variables in microleakage experiments

Radioisotope (50'ies) e.g.
⁴⁵Ca
¹³¹I
³⁵S
²²Na
¹⁴C
³²P
⁸⁶Rb

Neutron activation (70'ies)
²⁴Mg → ²⁶Mg
 (contaminants)

Dye penetration/extraction, e.g.
 Aniline dye
 Basic fuchsin
 Crystal violet blue
 Eosin
 Methylene Blue

(Fluorescent) penetration/extraction, e.g.
 Sodium fluorescein
 Acriflavin
 Auramine O
 Rhodamine-B

Specimen variables
 Tooth specimens & storage
 Cavity dimension/C-factor & location (dent. permeability)
 Margin location & isolation
Aging (Pre- OR in Immersion)
 Thermal cycling /°C /time / #
 Mechanical loading N / #
 pH cycling
 Food-simulating solutions

Other methods
 Air- / liquid pressure
 Bacteria
 Electrochemical
 Fluid filtration
 Moisture/«Percolation»

Chemical tracers
 AgNO₃ (“nano-leakage”) → hydroquinone developer

µCT

Dye variables
 Concentrations & pH
 Diameter/molecular weight
 Tissue /material affinity
 pH resistance
 Time
 Rinsing v/z contamination
 Sectioning/extraction 2D/3D
 Identification & resolution
 Quantification & analysis

Examples of variables in microleakage experiments

2% Rhodamine-B, 24h + Vacuum (Patil ea 2015)

0.5% Basic Fuchsin, 24h (Cehreli ea 2013)

silver nitrate 50% w/v, 24 h. (Rengo ea 2012)

1% methylene blue, 24 h. (Patil ea 2015)

0.5% methylene blue, 24 h. (Dalli ea 2013)

2% methylene blue, 48 h. (Arisu ea 2009)

0.5% Basic Fuchsin, 16 h. (Tantbirojn ea, 2011)

silver nitrate 50% w/v, 2 h. (Chandurkar ea 2012)

2% methylene blue, 48 h. (Arisu ea 2009)

Reviews on microleakage studies over last 4 decades

Author	Main conclusion
1968, Roydhouse	...of limited value ...because many variables are not accounted for. Tests may demonstrate a potential, but not a clinical reality
1969, Lelabelle et al.	...these tests eliminate the effect of pulpal hydrostatic pressure and plaque
1972, Going	...most methods fall scientifically short in providing quantitative data
1981, Jodaikin	...no direct comparison possible between in vitro & vivo due to many variables
1982, Shortall	...the results can be partly or totally influenced by the variations of the methodology applied
1992, Taylor & Lynch	...wide variations in methodologies are revealed
1991, Söderholm	...the relevance in a testing protocol for dentin adhesion must be questioned
2001, Raskin et al.	...results from different testing institutes could not be compared
2003, Raskin et al.	...results from different testing institutes are hardly reproducible
2007, Heintze	...it does not make sense to use this elaborate labor-intensive method
2007, Sarrett	...evidence for a direct relationship between poor marginal quality as promoter or primary cause for secondary caries is unlikely
2011, Schmid-Schwab et al.	...not possible to make a quantitative synthesis due to study heterogeneity
2011 Heintze & Zimmer	...dye penetration...do not correlate or correlate only partially with clinical findings
2012, Dennison & Sarr	...clinical evidence refute earlier conclusions that clinical microleakage leads to secondary caries
2012, Bayne	...no correlation of microleakage with any clinical event has ever been established
2013, Dietschi et al.	...the further use of this test method in the future should be strictly limited
2013, Heintze	...moderate evidence that dye penetration tests does not correlate with clinical data

Microleakage observations versus clinical observations

Clinical variable	Microleakage	Clinic
Incremental vs bulk filling	Less microleakage	Corroborate
Different curing approaches	conflicting results	?
Enamel vs dentin margins	Less microleakage	Corroborate
Light cured vs self-cured	Less microleakage	Corroborate
Matrix system	conflicting results	?
Primer solvent	Effect on dentin, not enamel	Corroborate
Incorrect cavity drying	More microleakage	Corroborate
Boxed cavity form	More microleakage than if rounded	Corroborate
Sharp margins	More microleakage than if beveled	?
Occlusal loading	More microleakage than if no loading	?
Thick flowable liner	Less microleakage in enamel (- dentin)	?
Adhesive brand	conflicting results	?
Flowable u. packable resin	conflicting results	?
rmGIC u. composite	conflicting results	?
Flowable u. rmGIC	conflicting results	?
Composite vs packable resin	Less microleakage in dentin	?
Etch-and-rinse vs self-etch	conflicting results	Corroborate
Single versus two layers	Less microleakage	Corroborate
Composite brand	conflicting results	?
Composite vs ormocer	conflicting results	?
Composite direct vs indirect	conflicting results	?

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- Secondary caries incidence in controlled clinical studies versus cross-sectional examinations

In-vitro caries-like lesions adjacent to restorations – Pioneer studies

Acidified broth

Mortensen ea. J Dent Res 1965

Acidified medium*

*Gelatin adjusted to pH=4 by addition of lactic acid, imbibed 5 - 200 days

* Silverstone (1968)

First used by Hals & Nemeas, Caries Res 1971.

Ellis & Brown. J Dent Res 1967 & Atto et al. J Dent Res 1970

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*Gelatin adjusted to pH=4 by addition of lactic acid, imbibed 5 - 200 days

* Silverstone (1968)

First used by Hals & Nemeas, Caries Res 1971, + ~30 papers, by e.g., Hals, Grieve, Kidd, Heintze, Zimmerman, Hicks, Donly and others)

Gelatin / gelatin gel or agar / methylcellulose

Ellis & Brown. J Dent Res 1967 & Atto et al. J Dent Res 1970

Cycling pH 4.7 ↔ 7

Common methodologies (Str. Mutans):
Dummer et al 1982
Fontana et al. 1996

«Lesions» along restorations detected in PLM – what was actually observed?

Hals & Nernaes, 1971:

- Usually ... a narrow subsurface defect ... gradually encompassed the whole filling without penetrating deeply into the tissue*
- Wall lesions were seen only when imbibed in quinoline and not in air or water & only in the PLM

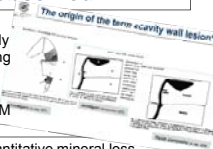
PLM is purely qualitative and does not provide quantitative mineral loss

Enamel

- Quinoline facilitates identification of caries due to pore size selectivity
- Penetration of ions into dental tissues change optical birefringence, e.g. corrosion products from amalgam such as oxides, sulphides and chlorides of tin, and to a lesser extent zinc and copper

Dentin

- Quinoline binds because of a «von Ebener phenol reaction» – i.e., a selective binding to collagen, and not due to pore penetration
- Dentin tubules demonstrates form birefringence due to their micrometer size
- Collagen displays form birefringence



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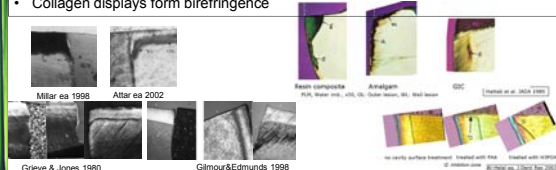
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Effects of the acidified procedure on the restoration-tooth interface?

- Several strategies - often without consideration of likely negative effects on the restorative material and not occurring in reality intra-orally
- Restored teeth sometimes exposed directly after the setting time, i.e., not always synonymous with a fully hardened or polymerized material
- In aggressive media crevice corrosion cells are likely generated in the interface along metallic restorations, which lowers the pH further, and cements such as glass ionomers undergo profound surface erosion
- The adoption of methodologies for causing artificial caries-like lesions in enamel were perhaps too uncritically extrapolated to create artificial caries-like lesions adjacent to restorations

More common procedure today:
pH cycling: 4.4 ↔ 7 **

** Featherstone et al. 1983/86



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- In aggressive media crevice corrosion cells are likely generated in the interface along metallic restorations, which lowers the pH further, and cements such as glass ionomers undergo profound surface erosion
- The adoption of methodologies for causing artificial caries-like lesions in enamel were perhaps too uncritically extrapolated to create artificial caries-like lesions adjacent to restorations (Featherstone, 1996)
- Dental caries is not limited only to demineralization, but becomes heavily infected by mono- or multispecies biofilms, which is difficult to reproduce fully in vitro
- Confounder when the research focus is demineralization-reminerzalization of artificial caries-like lesions adjacent to restorations made from materials with alleged anticariogenic properties

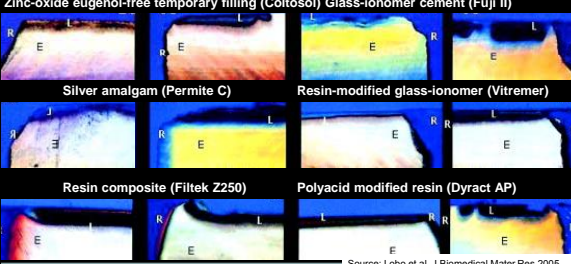
Effects of the acidified procedure on the restoration-tooth interface?

Sa et al. 2004: *Glass ionomers demonstrated significant anti-cariogenic properties when exposed to the chemical model. However, no significant anti-cariogenic properties were observed with the microbial caries model. In conclusion, ...caution should be exercised when trying to extrapolate the results of in vitro studies to the clinical situation*

Zinc-oxide eugenol-free temporary filling (Coltisol) Glass-ionomer cement (Fuji II)

Silver amalgam (Permite C) Resin-modified glass-ionomer (Vitremer)

Resin composite (Filtek Z250) Polyacid modified resin (Dyract AP)



Source: Lobo et al. J Biomedical Mater Res 2005

Topics discussed in this presentation

- Dental caries, a brief review
- Restorative materials and the tooth-restoration interface
- The (cavity) “wall lesion” – what is in a word?
- Etiopathogenesis of secondary caries gained from in vitro research
 - Microleakage
 - Artificial caries-like lesions adjacent to restorations
- Secondary caries incidence in controlled clinical studies versus cross-sectional examinations

The replacement of dental restorations

Primary reason identified in:
1991: Secondary caries - 2001: Secondary caries - 2012: Secondary caries

Longevity of posterior restorations

L. A. Miles, A. J. Johnson and V. G. Dale

Quality of dental restorations
FDI Commission Project 2-95*

Longevity of posterior composite restorations:
Not only a matter of materials!

Operator
Patient
Material

Based on compilations of:

- Observational data from:
 - Cross-sectional studies of reasons for replacement of restorations, occasionally with true or estimated time since placement
 - Cross-sectional studies of restorations in situ, occasionally with true or estimated time since placement
 - Prospective & retrospective studies of patient cohorts or subgroup analyses of such
- Experimental studies, with variable internal or external validity reflected by study power, randomization, likelihood of confounding & risk of biases

Which estimates should we trust?

- Estimates of incidence & prevalence of secondary caries have ranged from insignificant to extensive. Scepticism have been voiced in both directions
- Potential biases that likely influences estimates is extensive
 - selection bias - performance bias - detection or assessment bias - attrition bias - reporting bias
- Typical examples: patient recruited amongst dental students and faculty; studies not conducted amongst GPs, lack of operational descriptive criteria or judgement of own clinical work; high number of patient dropouts especially amongst the unhappy ones; and the reporting of surrogate outcomes rather than patient-relevant ones
- Results based on clinical work in settings where cost per unit time is of nominal concern do not provide any indications on how the restorative material will perform when placed by the average dentists in mouths of their spectrum of patients during a busy workday.
- The data sampling method, patient demography as well as study methodology influence estimates - Is a quest for "overall" exact values meaningful from a scientific or clinical perspective?

When the «science» in evaluating «scientific research» becomes an exercise in nihilism

Longevity of posterior resin composite restorations in adults - A systematic review

Alfacher Amaladoss^{1,2}, Jorana Daghfles³, Jon W. H. van't Hof⁴, Aron Nouri Akbar⁵, Candice Sandberg-Englund⁶, Sofia Thonness^{7,8}, Mikael Mattsson⁹

Fig. 1 - Selection process for study inclusion.

i.e.,
RCTs > 4 yrs &
<5% attrition/year

Only 6 / 4275 papers = 0,0019% was considered for grading of scientific evidence and conclusions

- In today's mobile world, the likelihood of a near-zero attrition is unrealistic.
- At what level does the attrition rate in a dental study become a concern with regard to restoration performance estimates?
- Good research ethics allow study participants at any time to drop out without having to explain why. Coercive offers is generally regarded as unethical.
- Are study participants who return for a follow-up clinical examination many years later in the same clinic representative of the general population when it comes to oral health attitudes and treatment behavior?

What are the alternatives for collecting data?

Health register data analyzed with Multi-level regression statistics (AKA hierarchical linear r., nested models r., mixed models r., random coefficient r., random-effects models r., random parameter models r.)

- UK: Burke & Lucarotti (80K+ adults)
- Finland: Vähänikkilä / Käkilehto / Suni (6K, 36K adults)
- USA: Bogacki / Coppola (300K / 1.500K adults)
- Norway: Dobloug & Grytten (64K adults)
- Brazil: Demarco & Correa (6K adolescents)

Pragmatic (real-life) studies, e.g., in Practice-based Research Networks

- Data on restoration performance obtained in cross-sectional studies reflect the good and the bad operators and oral health attitudes and practices of patients.
- However, how many dentists in real-life
 - prepare textbook-like cavities in teeth to receive the restoration?
 - handle and place restorative materials according to handling instructions?
 - ensure that their patients are motivated and enabled to prevent future caries?

CONCLUSIONS

Conclusions - 1/3 "caries wall lesions"

- It is doubtful whether caries can exist in the restoration-tooth interface independently of an outer enamel caries lesion.
- The term "wall lesion" including its variants is ill defined, has been, and is still being used indiscriminately.
- Stakeholders should avoid using this ambiguous label due to its connotation to an entity that does not exist per se.

Conclusions - 2/3, experimental data

- Microleakage experiments continue to emerge regardless of multiple reviews questioning the reliability and validity of the method.
- Several of the approaches used to generate artificial caries-like lesions are very aggressive. Remarkably little discussion has evolved about how these aggressive approaches create microenvironments that do not occur in reality. Corrosion- and biodegradation products may influence the biofilm qualitatively and quantitatively and it is difficult to replicate these variables in any ex vivo environment.

Conclusions - 3/3, clinical data

- Clinical data sampling method, patient demography as well as study methodology influences the incidence and prevalence estimates of secondary caries.
- Clinical results based on clinical work in settings where cost per unit time is of nominal concern do not provide any indications on how the restorative material will perform when placed by the average dentists in the mouths of their spectrum of patients during a busy workday.

One of the most gorgeous sites on Maui to watch sunrise & sunset: The Haleakala mountain:

If you plan to see the sunset on top of Haleakala mountain - An important safety message:

**AT:
6.13 PM
40 °F**

If you plan to see the sunset on top of Haleakala mountain - An important safety message:

**AT:
6.18 PM
40 °F**

SONIC FROM HERTZ WITH A SHORT-CIRCUITED REMOTE CONTROL

If you plan to see the sunset on top of Haleakala mountain - An important safety message:

**AT:
6.28 PM
35 °F**

SONIC FROM HERTZ WITH A SHORT-CIRCUITED REMOTE CONTROL

What are your options?



1. Panic
2. If you have a satellite telephone:-Call the Hertz emergency centre in Florida and remember how to spell H-a-l-e-a-k-a-l-a before they can help you
3. Start walking down the 10.023 feet mountain in the dark
4. Hijack one of the 4 remaining cars on the parking lot or meet a good Samaritan

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5. Remove the battery in your remote controller – (non-validated test)

