



# A review of the questions and needs in endodontic diagnosis

I. ABU-TAHUN, M.A. AL RABAB'AH, A. KHRAISAT

*Dpt of conservative dentistry and fixed prosthodontics, Faculty of Dentistry, The University of Jordan, Amman, Jordan.*

## Abstract

The current diversity of opinions in endodontic diagnosis has been a source of interest and academic debate by clinicians and researchers. Currently, no single pulp testing technique can reliably diagnose all pulpal conditions neither it has been proven to be superior in all aspects.

Despite improvements of various aspects of this process, there are no historically dramatic changes, or consensus for pulpal status in health or disease in addition to a lack of relative systematic reviews.

In this review, the past, present and future most debated and critically questioned issues of endodontic diagnosis are discussed. The aim of this review is to provide insights in future diagnostic modalities and areas for further study in endodontic practice pertinent to diagnosis.

**Keywords :**  
**Diagnosis,**  
**pulp testing,**  
**endodontics**

## Résumé

### Revue des questions et besoins dans le diagnostic endodontique

La diversité actuelle des opinions en endodontie diagnostique a été une source d'intérêt et de débat académique chez les cliniciens et chercheurs. Actuellement, aucune technique de test de vitalité pulpaire ne peut diagnostiquer avec fiabilité tous les états pulpaire ni prouver sa supériorité.

Malgré les progrès des divers aspects de ce procédé, il n'y a pas de changement historiquement notable ou consensus sur l'état de la pulpe saine ou malade, avec de plus un manque d'études relatives systématiques.

Dans cette étude, le passé, le présent et le futur des questions les plus débattues sur le diagnostic endodontique sont discutées. Le but de cette étude est donner un aperçu des futures modalités et domaines diagnostiques pour de prochaines études de pratique endodontique utiles au diagnostic.

**Mots-clés :**  
**Diagnostic,**  
**test pulpaire,**  
**endodontie**

## Introduction

Diagnosis, is "the art and science of detecting deviations from health and the cause and nature thereof" where the data obtained from questioning, examining and testing are combined by the dentist to identify deviations from the normal (1, 2).

Aim of endodontic diagnosis is to prevent irreversible pulpal injuries and apical periodontitis and thereby optimize the outcomes of preventive and interventional endodontic treatments (3).

All methods to determine the status of the pulp and root-supporting structures except for

surgical exploration and histological examination (biopsy), rely on indirect diagnostic data interpreted from the patient response to a stimulus placed externally to the tissue (4, 5).

Due to this inability to directly test the pulp, testing results are based on assumptions of what is presumed to be the underlying disease process of a given clinical state with multiple clinicians arriving at vastly different interpretations of the same data (6).

Tests used with "yes" or "no" response which varies from patient to patient can generally identify patients free of disease but are less

effective in identifying patients who have pulp disease (7).

Extra testing measures advised to substantiate a more predictable result and attain a reproducible reading, although predictive in some cases, were not able to solve the problem of indirect response of the patient.

Literature on diagnosis of pulpal status is almost devoid in the area of permanent teeth with immature apices particularly following trauma (6, 8).

The aim of this comprehensive search of the literature from September 1954 to May 2010 was to critically discuss the different grey areas in actual diagnostic process to address challenges and needs and to plan a more evidence based thinking for researchers and academics to enhance clinical care particularly in treating developing teeth.

## Materials and methods

A library electronic search of MEDLINE, PubMed, and Cochrane databases using specific MeSH terms from September 1954 to May 2010 was made. Exhaustive hand searching and citation mining for all relevant articles as well as classic and meaningful endodontic textbooks reporting endodontic diagnostic tests and their relevance to clinical situations in traumatized and developing teeth was also targeted.

## Study selection

Following this literature search, the titles and abstracts of all articles identified from the electronic and hand searches were first screened to select articles that clearly meet the search criteria and selected articles were reviewed to develop consensus that the inclusion and exclusion criteria were respected.

## The diagnostic dilemma

Clinical pulpal states of health and disease we use today were early described by MORSE et al. (9), but there is little or no correlation between clinical diagnostic findings and the histopathologic state of the pulp (10, 11).

Different classification systems have been advocated for pulp diseases based on histologic findings or mixing clinical and histologic terms (12, 13). LINDA et al. 2009 (14), identified 14 different classification systems for pulp diseases over the years most of them are based on histologic findings.

The result was a misleading diagnosis for the same clinical condition creating confusion when a rational treatment plan needs to be established. To describe various disease states of the pulp, new clinical classification scheme were added with claims of superiority to enhance the accuracy and clinical relevance of diagnostic terminology (15-18).

Diagnosis is needed to perform clinical endodontic treatment, and histopathologic diagnosis is impractical in daily endodontic practice with questionable value of clinical detection of a histologic change as a diagnostic term in some cases (19-22). This perception of treatment which is not actually different with the different diagnostic terms raised the question as to whether diagnosis should be a separate activity from treatment (5).

Of all the histopathological pulpal states, the term pulp necrosis used to classify death of the pulp, which is, for the most part, a histological finding is successful in terms of the extent of possible canal infection and the use of the term pulp necrosis is most reliably predicted from a clinical testing point (11, 18).

While distinction between partial and full necrosis becomes important when dealing with immature teeth that have an open apex (13, 23), in exposed pulps of children, test results

and clinical symptoms do not coincide with pulpal histology complicating diagnosis in these teeth (24). The correlation between a clinical diagnosis and the histological status of the pulp as reported by GARFUNKEL et al. was about 50%. All 25 previously traumatized anterior teeth that reacted negatively to conventional pulp testing contained vital pulps when examined histologically (25).

### **Biological considerations**

Understanding the biology of the dental pulp and demarcation of pulpal inflammation and healing (26), is fundamental to the understanding of patient response to different testing modalities (27) and would result in more biologically and ethically oriented treatment options (28).

Diagnostic tools to determine the extent of pulpal inflammation are imprecise and studies that have attempted to determine diagnostic accuracy of reversible versus irreversible pulpitis are few (28). Clinical tests available can only test the ability of the pulp to respond to a stimulus (21) which does not represent our best clinical judgment for the actual state of the pulp where the coronal pulp might be infected but the apical pulp remains vital with a varying degree of inflammation (29).

Another area of debate is the presence of cellulitis or acute exacerbation that present a more complex etiological and therapeutic situation (30). Identification, quantification and link of bacterial and biologic markers and genetic assays with pulpal inflammation and symptomatic teeth will help in controlling endodontic infection and its complications and will affect the ultimate outcome of endodontic treatment (31, 32).

Based on the individuality of each lesion and the progression of pulp diseases through various stages, a comprehensive clinical diagnostic approach and steps in examination must be recognized. Since etiologies, treatment strategies, and prognoses vary considerably,

the term "differential diagnosis" was also suggested as more realistic in its application in endodontics, especially when tooth pain is the chief complaint (33, 34).

### **The pain system in the pulp**

The inconsistent definitions of pulpal disease have led many researchers to classify pulpal status into two main general categories: vital or nonvital or response versus no response (35).

The dental information gathering process and the history of periradicular pain for teeth with necrotic pulps, is an acceptable initial pain-assessment tool for endodontic emergency patients that might increase the accuracy of pain localization and aid in determining a pulpal diagnosis, but it would not yield predictive value for many patients (36-38).

Subjective symptoms are only partially related to the status of the pulp and are occasionally misleading (39). This relatively straightforward diagnostic outcome becomes more difficult to interpret particularly in posterior teeth (40) when the area would "ache all over" with increasing discomfort (41).

Although pain is strongly synonymous to endodontics, most endodontic pathoses are asymptomatic and pain cannot be used to differentiate endodontic problems from non endodontic pathoses (42, 43).

Distinguishing true pain originating from an irreversible pulpitis that may indicate the need for root canal treatment versus hypersensitivity that may indicate the need for palliative management is important (44,45) .

The considerable number of techniques described for pain measurement including verbal and numeric rating scales (46, 47), visual and color analog scales (48), finger span expression (49), calibrated questionnaires (50), cortical evoked potentials (51), and formatting the process using systematic format such as S.O.A.P., which is an acronym for Subjective findings, Objective tests, Assessment (or Appraisal), and Plan of treatment to increase efficiency and consis-

cy indicate the significant deficit of pulpal pain assessment (52).

Statistical analysis by GRUSHKA and SESSLE (53), using the McGill Pain Questionnaire to differentiate valid predictors of whether pulp inflammation is reversible or irreversible did not allow an acceptable level of determination accuracy. None of the other metrics such as the history of presenting symptoms (54) or a history of being spontaneous (10) could result in sensitivity, specificity, or positive/negative predictive value of the symptoms.

Subcategories of classification of pulpitis and systematic evaluation of pain that represent the biologic rationale for endodontic diagnostic tests are required to enable practitioners differential diagnosis of which pulps can be managed conservatively and which ones require more radical treatment including extraction of the tooth (14, 55).

As a result of the vague descriptors relative to the pain experienced by the patient, recent forms of more precise pain measurement including electroencephalography and standardizing the measurement of mechanical allodynia were introduced, yet, their value in endodontic diagnosis and treatment are to be determined (56-59).

#### **Diagnostic concerns in traumatized and developing teeth**

Many attempts have been made over the years to classify dental injuries. The currently accepted system applicable to injuries to the teeth and supporting structures that can be applied to both primary and permanent dentitions is based on the World Health Organization's Application of International Classification of Diseases to Dentistry and Stomatology, and its modification by ANDREASEN (60).

Even though definite diagnosis is established only after inspection and probing of the pulp chamber and the root canal, examination of a patient with dental injuries often includes

chief complaint, history of traumatic injury to the facial area, pertinent medical history, and clinical examination (61).

The value of electric pulp tester (EPT), currently the most used test to assess the neurovascular supply to the pulp of a traumatized tooth drops considerably following trauma in which conditions the innervation of the pulp might be jeopardized, at least temporarily (62) and may lead to false negative or false positive responses as roots mature (6, 63).

The less variability in findings for specificity and sensitivity of electric pulp tests renders them more consistent at identifying teeth without disease (vital pulp) (6, 29).

After a luxation injury in traumatized young teeth with wide-open apices in either developing or even mature teeth, lack of response to EPT should not automatically be accepted as proof of pulp necrosis (5, 64). On the other hand, the response at the time of first injury should also be interpreted with caution as sensory nerves may not yet have developed fully and the response might also be affected by overreaction of the child to the stimulus (34).

There is no agreement as to whether thermal tests, when used in the absence of other tests, can reliably determine the presence of a diseased pulp or to identify teeth without disease (29). These qualitative tests can only determine health versus disease caused by a particular primary afferent nerve response and, by necessity, the patient's symptoms (65). In teeth with open apices after traumatic injury, these tests may be unreliable, no response might be elicited even after circulation has been restored and heat tests in permanent teeth with developing apices are rarely performed (8).

Ice and ethyl chloride are of limited value and have consistently been reported to be inferior to carbon dioxide snow shown reliable by many studies concerned with vitality determinations in luxated, avulsed, or root fractured

teeth (54, 62, 63, 66) (8, 67), and dichlorodifluoromethane (DDM) (68).

The oldest pulp vitality tests, palpation and percussion, may be reliable in identifying inflammation in the periodontal ligament space and can also provide information about the relationship between the tooth and adjacent bone indicating lateral or intrusive displacement (69), but cannot differentiate pulpal from periodontal diseases (70).

A positive response to the biting stress test is highly suggestive of periodontal inflammation or incomplete crown-root fracture (59). Mobility and periodontal pocket depth are more standardized than percussion, palpation, and biting stress tests. Even though type of luxation can be related to the degree of mobility, information regarding changes in the root-supporting structures is limited (60).

Transient coronal discoloration has been reported in 4% of teeth after luxation injuries as a result of transient apical breakdown after displacement injuries (26, 71). Transient periapical radiolucency, together with coronal discoloration, negative electric pulp test, and cold response up to 4 months, was shown to subsequently regain the original color and normal pulpal responses when healing is complete. To avoid mistakes, there should be no rush in treatment undertaken on the basis of negative responses (26, 71).

#### **Do radiographs tell the truth ?**

A routine radiograph does not reveal the third dimension, which is important in teeth with an open apex. In addition, correlation between radiographic and histologic diagnosis is poor.

Showing better specificity than sensitivity, conventional radiographs are better able to identify the teeth without periapical disease than to identify the teeth that have periapical disease (72, 73), leading to unreliable inter and intraexaminer agreement on interpretation of structural changes in the periradicular

tissues (64, 74).

Following traumatic injuries, two or three periapical x-rays taken from different angles were suggested to increase the accuracy of the radiographic interpretation of the changes in the dentoalveolar complex (64, 75, 76).

A relatively high probability of a false-negative result with both periapical and panoramic imaging techniques was reported preventing reliable differentiation of periapical cysts and granulomas made with conventional periapical radiographs (77).

Another difficulty is encountered when radiographic observation is used as predictable criterion for revascularization and continued root development. The difficulty lies in obtaining the same film position, and the possibility to distinguish arrested root formation and complete development (78).

#### **On the horizon**

Radiographic improvements have reduced radiation exposure and improved convenience visualization of changes in a measurable way. Digital and digital subtraction radiography appears to enhance and improve the ability to detect and measure the size of periradicular lesions and may improve diagnostic accuracy particularly in the evaluation of healing (79, 80).

The potential of Ultrasound imaging introduced in endodontics by COTTI et al (81) agreed with the histopathological diagnosis in all 15 cases examined. The wide spectrum applications of cone beam volumetric tomography (CBCT) in endodontics apart from evaluating endodontic treatment outcomes include diagnosis, detection of canal morphology, non endodontic pathosis, root fracture, internal resorption, invasive cervical resorption, anatomic presurgical assessment (82,83).

As (CBCT) machines become more common in dental offices, CBCT may be the answer to more early and accurate diagnosis of peri-apical

pathosis and may resolve the issue of inter- and intraobserver interpretation of radiographic images (5). Despite the slight variation in sensitivity due to tooth location, very high specificity was found in all tooth types for both imaging techniques (77).

While for teeth with fully formed roots, clinical diagnostic determinations for endodontic therapy depend on whether the pulp spaces are infected, in vital pulp therapy to preserve the pulp of teeth with incompletely formed roots, primary question is whether the treated pulp remains healthy (84, 85).

Developing teeth have the potential of regeneration and revascularization of the injured pulp after trauma and having information about the pulp status of traumatized teeth can be of great value. Measurement of tooth surface temperatures was widely used as a step ahead to determine pulp vitality. A review of the literature reported that this technology is not sensitive enough to identify periapical lesions deep within the bone that would indicate a necrotic pulp or irreversible pulpitis (86). The usefulness of the technique in endodontics needs improvement of science and technology (87).

Several experimental methods have been used to assess pulpal blood flow. These include invasive methods such as radioisotope clearance (88), H<sub>2</sub> gas desaturation (89), and non invasive techniques such as Laser Doppler Flowmetry (LDF) (90), pulse oximetry, photoplethysmography, and dual wave length spectrophotometry (91).

Transmitted light photoplethysmography (TLP), is a non-invasive technique successfully used to monitor pulpal blood flow in animal and human studies causing less signal contamination from the periodontal blood flow than is the case for LDF (92).

LDF, initially designed in the early 1970s to measure blood flow in the retina and pulse oximetry are two gold standards of vitality tests, having higher sensitivity and specificity than cold, heat, and electric tests (93, 94).

The only few investigations of pulpal vitality using this approach indicated that the consistency of time between peaks in pulses would give an indication of vitality in a tooth pulp by establishing pulsatile reading of similar frequency to the heart rate and might prove a more feasible method to be used in dentistry (95, 96).

The use of LDF in dental trauma has proven to be more valuable than in assessing vitality in healthy pulps especially those with history of trauma or luxation injuries where LDF can detect revascularization after a few weeks, and well in advance of other more traditional clinical tests (97, 98).

Showing signs of adverse outcomes in luxated teeth, LDF may provide opportunity to identify "at-risk" teeth early after the trauma, and initiate treatment with confidence prior to the tooth being lost from pulpal necrosis and infection (99, 100).

The technology of pulse oximetry has allowed significant advances in the medical field.

Approximately 25 models of the pulp oximeter are available to provide pulse rate and oxygen saturation readings calculated in a micro-processor (101). The distinct advantage offered by this pulp testing method in trauma cases will allow unique opportunity for an immediate objective diagnosis of vascular integrity and diagnosis of the pulpal vitality (102).

The accuracy rate of detecting negative test result to indicate a vital pulp by the use of a custom-made pulse oximeter probe specifically made for dental application compared with thermal and electric pulp tests, was found by GOPIKRISHNA and coll. to be 74% with the electrical tests, 81% with the cold test, and 100% with pulse oximetry (62).

The accuracy of this diagnostic method, the completely noninvasive nature and superior patient acceptance support the need for additional studies in the use of the pulse oximeter to interpret the pathological processes of the



pulp (102-104).

Advances in measuring pulpal blood flow still in their infancy and the high-resolution, 3-dimensional imaging may allow better correlation between pulpal histopathologic states and clinical phenomena (14, 105, 106). Used within their limitations, all findings of these technologies are promising in assessing pulpal vitality in healthy and traumatized teeth (98).

Future developments could possibly be a part of using easier, less costly and more refined oral diagnostic procedures that may have a better chance of success. The use of specific biochemical markers in the gingival crevicular fluid such as electrophoresis technique proposed in the early 1970s to differentiate between periapical cysts and granulomas, may yield future useful diagnostic tools (107, 108, 109).

### Conclusions and recommendations

Despite that current (old) diagnostic tests still hold a place of respect results from the literature suggest that endodontic practitioners are supportive and optimistic about the future use of more sophisticated and noble endodontic procedures (110). There is a gap in the evidence and a gap in the knowledge to support and validate what we are doing in the actual diagnostic practice, yet, exploration of new testing devices, approaches and materials in the new era of endodontic practice appear to have improved in theory and application giving a better picture as to what the dental pulp might appear as histologically (111-113).

Because of the relatively few evidence-based, randomized, controlled clinical trials in this topic area, in addition to the lack of treatment-oriented diagnosis scheme, a more reliable body of scientific evidence is the more pressing need today to find out an ideal metric, or combination of metrics, that would result in greater specificity and sensitivity and higher levels of evidence to select the appropriate treatment modalities (19).

In addition to being inexpensive, any diagnostic method to arise in the near future should be able to discriminate more accurate pulpal conditions tested than today, based on the best available evidence to produce predictive value for pulpal pathology in a clinical setting. On the basis of available evidence, it appears that better and more accurate quantification modalities of periradicular pain may lie in devices that allow direct measurements of pain thresholds as well (19). A new clinical-related classification which will improve communication among clinicians and researchers and unify practitioners is essential. This new classification should be simple to learn and teach, and should include the most common types to develop universally accepted criteria. The arbitrary use of terms, without taking into account the historic basis for the endodontic diagnostic scheme, may very well lead to overtreatment.

The importance of diagnostic skills in the practice of endodontics has been underscored by a recent 2008 AAE-sponsored symposium on endodontic diagnosis (66). Diagnostic process is not pure science, and the necessary examining equipment may not be the diagnostic tool or instrument but the diagnostician who will perform the test and arrive at a reliable conclusion. Practitioners should be equipped with the basic requirements and skills including the art of listening, knowledge, training, interest, curiosity, patience and above all common sense. As educators and instructors, it's our responsibility and duty to declare that many aspects of the true "puzzlement" could be attributed to the scope of university training, as diagnostic skills might be beyond the comfort level of the students.

Having an influence on subsequent endodontic decision making and treatment, referral should be considered, particularly if diagnosis reaches a dead end or the chief complaint is not of endodontic origin.

## References

1. **GIFT HC, BHAT M.** Dental visits for orofacial injury: defining the dentist's role. *J Am Dent Assoc* 1993 ; 124 : 92-8.
2. **AMERICAN ASSOCIATION OF ENDODONTISTS** Glossary of endodontic terms. 7th ed. Chicago: American Association of Endodontists ; 2003.
3. **CAMP JH.** Diagnosis dilemmas in vital pulp therapy : treatment for the toothache is changing, especially in young, immature teeth. *J Endod* 2008 ; 34 (Suppl 7) : 6-12.
4. **ROSENBERG PA, SCHINDLER WG, KRELL KV, HICKS ML, DAVIS SB.** Identify the endodontic treatment modalities. *J Endod* 2009 ; 12 : 1675-94.
5. **NEWTON CW, HOEN MM, GOODIS HE, JOHNSON BR, MCCLANAHAN SB.** Identify and determine the metrics, hierarchy, and predictive value of all the parameters and/or methods used during endodontic diagnosis. *J Endod* 2009 ; 12 : 1635-44.
6. **FUSS Z, TROWBRIDGE H. IB, et al.** Assessment of reliability of electrical and thermal pulp testing agents. *J Endod* 1986 ; 12 : 301-5.
7. **REIT C, KVIST T.** Endodontic retreatment behaviour r: the influence of disease concepts and personal values. *Int Endod J* 1998 ; 31 : 358-63.
8. **FULLING HJ, ANDREASEN JO.** Influence of maturation status and tooth type of permanent teeth upon electrometric and thermal pulp testing procedures. *Scand J Dent Res* 1976 ; 81 : 286-90.
9. **MORSE DR, SELTZER S, SINAI I, BIRON G.** Endodontic classification. *J Am Dent Assoc* 1977 ; 94 : 685-9.
10. **SELTZER S, BENDER IB, ZIONTZ M.** The dynamics of pulp inflammation : correlations between diagnostic data and actual histologic findings in the pulp (part I). *Oral Surg Oral Med Oral Pathol* 1963 ; 16 : 846-71.
11. **SELTZER S, BENDER IB, ZIONTZ M.** The dynamics of pulp inflammation: correlations between diagnostic data and actual histologic findings in the pulp (part II). *Oral Surg Oral Med Oral Pathol* 1963 ; 16 : 969-77.
12. **ABBOTT PV.** Classification, diagnosis and clinical manifestations of apical periodontitis. *Endod Topics* 2004 ; 8 : 36-54.
13. **ABBOTT PV, YU C.** A clinical classification of the status of the pulp and the root canal system. *Austral Dent J* 2007 ; 52 (Suppl) : S17-31.
14. **L.G. LEVIN, A.S. LAW, HOLLAND, C. ENDO, PAUL V. ABBOTT, R.S. RODA.** Identify and define all diagnostic terms for pulpal health and disease states. *J Endod* 2009 ; 35 : 1645-1657.
15. **GLICKMAN GN, MICKEL AK, LEVIN LG, FOUAD AF, JOHNSON WT.** Glossary of endodontic terms. 7th ed. Chicago : American Association of Endodontists. 2003.
16. **AMERICAN BOARD OF ENDODONTICS.** Pulpal & periapical diagnostic terminology. Chicago: American Board of Endodontics ; 2007.
17. **ABBOTT PV.** The periapical space : a dynamic interface. *Ann R Australas Coll Dent Surg* 2000 ; 15 : 223-34.
18. **NECROSIS.** Dictionary.com. Merriam-Webster's medical dictionary. Merriam-Webster, Inc. Available at: <http://dictionary.reference.com/browse/necrosis>.
19. **A.L. FRANK, M. TORABINEJAD.** Diagnosis and treatment of extracanal invasive resorption. *J Endod* 1998 ; 24 (7) : 500-4.
20. **STANLEY HR, PEREIRA JC, SPIEGEL E, BROOM C, SCHULTZ M.** The detection and prevalence of reactive and physiologic sclerotic dentin, reparative dentin and dead tracts beneath various types of dental lesions according to tooth surface and age. *J Oral Pathol* 1983 ; 12 : 257-89.
21. **PASHLEY DH, LIEWEHR FR.** Structure and functions of the dentin-pulp complex. In : Cohen S, Hargreaves KM, eds. *Pathways of the pulp*. 9th ed. St Louis: Mosby-Elsevier; 2006. p. 460-513.
22. **NEWTON CW, COIL JM.** Geriatric endodontics. In: Cohen S, Hargreaves KM, eds. *Pathways of the pulp*. 9th ed. St Louis : Mosby-Elsevier ; 2006. P. 883-917.
23. **SMULSON MH, SIERASKI SM.** Histophysiology and diseases of the dental pulp. In : Weine FS, ed. *Endodontic therapy*. 5th ed. St Louis: Mosby ; 1996. P. 84-165.
24. **MASS E, ZILBERMAN U, FUKS AB.** Partial pulpotomy: another treatment option for cariously exposed permanent molars. *J Dent Child* 1995 ; 62 : 342-5.
25. **GARFUNKEL A, SELA J., ULMANSKY M.** Dental pulp pathosis. Clinicopathological correlations based on 109 cases. *Oral Surg Oral Med Oral Pathol* 1973 ; 35 (1) : 110-7.
26. **ANDREASEN F.** Transient apical breakdown and its relation to color and sensibility changes. *Endod Dent Traumatol* 1986 ; 2 : 9-19.
27. **JOE H.** Camp diagnosis dilemmas in vital pulp therapy: treatment for the toothache is changing, especially in young, immature teeth. *J Endod* 2008 ; 34 : S6-S12.
28. **GARFUNKEL A, SELA J, ULMANSKY M.** Dental pulp pathosis ; clinicopathological correlations based on 109 cases. *Oral Surg* 1973 ; 35 : 110-4.
29. **HYMAN JI, COHEN ME.** The predictive value of endodontic diagnostic tests. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 1984 ; 58 (3) : 343-6.
30. **P.N.R NAIR.** On the causes of persistent apical periodontitis: a review. *Int Endod* 2008 ; 39 : 249-81.
31. **JACINTO RC, GOMES BPPA, SHAH HN, et al.** Quantification of endotoxins in necrotic root canals from symptomatic and asymptomatic teeth. *J Med Micro* 2005 ; 54 : 777-83.
32. **SLOTS J, NOWZARI H, SABETI M.** Cytomegalovirus infection in symptomatic periapical pathosis. *Int Endod J* 2004 ; 37 : 519-24.
33. **BERGENHOLTZ G.** Effects of bacterial products on the inflammatory reactions in the dental pulp. *Scand J Dent Res* 1977 ; 85 : 122-129.
34. **ANDREASEN JO, ANDREASEN FM, BAKLAND LK, FLORES MT.** Examination and diagnosis. In: *Traumatic Dental Injuries: A Manual, 2nd edn.* Oxford: Blackwell Munksgaard ; 2003.p. 18-21.
35. **PETERS DD, BAUMGARTNER JC, LORTON L.** Adult pulp diagnosis: I - evaluation of the positive and negative responses to cold and electrical pulp tests. *J Endod* 1994 ; 20 : 506-11.
36. **MCCARTHY PJ, MCCLANAHAN S, HODGES J, BOWLES WR.** Frequency of localization of the painful tooth by patients presenting for an endodontic emergency. *J Endod* 2010 ; 36 : 801-5.
37. **BARBAKOW FH, CLEATON-JONES P, FRIEDMAN D.** An evaluation of 566 cases of root canal therapy in general dental practice: 2-postoperative observations. *J Endod* 1980 ; 6 : 485-9.
38. **BROWN B.** The measurement of human dental intrapulpal pressure and its response to clinical variables. *Oral Surg Oral Med Oral Pathol* 1965 ; 19 : 655-8.
39. **LIPTON JA, SHIP JA, LARACH-ROBINSON D.** Estimated prevalence and distribution of reported orofacial pain in the United States. *J Am Dent Assoc.* 1993 ; 124 : 115-21.
40. **FRIEND LA, GLENWRIGHT HD.** An experimental investigation into the localization of pain from the dental pulp. *Oral Surg Oral Med Oral Pathol* 1968 ; 25 : 765-74.
41. **VAN HASSEL HJ, HARRINGTON GW.** Localization of pulpal sensation. *Oral Surg Oral Med Oral Pathol* 1969 ; 28 : 753-60.
42. **NAHRI M.** The characteristics of intradental sensory units and their responses to stimulation. *J Dent Res* 1985 ; 64 : 564-571.
43. **P. MASCIA, B.R. BROWN, S. FRIEDMAN.** Toothache of non odontogenic origin: a case report. *J Endod* 2003 ; 29 : 608-10.
44. **LINSUWANONT P, PALAMARA JE, MESSER HH.** Thermal transfer in extracted incisors during thermal sensitivity testing. *Int Endod J* 2008 ; 41 : 204-10.
45. **MILLER SO, JOHNSON JD, ALLEMANG JD, et al.** Cold testing through full-coverage restorations. *J Endod* 2004 ; 30 : 695-700.
46. **GANGAROSASR LP, CIARLONE AE, NEAVERTH EJ, JOHNSTON CA, SNOWDEN JD, THOMPSON WO.** Use of verbal descriptors, thermal scores and electrical pulp testing as predictors of tooth pain before and after application of



• A review of questions... •

benzocaine gels into cavities of teeth with pulpitis. *Anesth Prog* 1989 ; 36 : 272-5.

47. **OWATZ CB, KHAN AA, SCHINDLERWG, SCHWARTZ SA, KEISER K, HARGREAVES KM.** The incidence of mechanical allodynia in patients with irreversible pulpitis. *J Endod* 2007 ; 33 : 552-6.
48. **MCCONAHAY T, BRYSON M, BULLOCH B.** Clinically significant changes in acute pain in a pediatric ED using the Color Analog Scale. *Am J Emerg Med* 2007 ; 25 : 739-42.
49. **FRANZEN OG, AHLQUIST ML.** The intensive aspect of information processing in the intradental A delta system in man: a psychophysiological analysis of sharp dental pain. *Behav Brain Res* 1989 ; 33 : 1-11.
50. **FALACE DA, REID K, RAYENS MK.** The influence of deep (odontogenic) pain intensity, quality, and duration on the incidence and characteristics of referred orofacial pain. *J Orofac Pain* 1996 ; 10 : 232-9.
51. **MOTOHASHI K, UMINO M, FUJII Y.** An experimental system for a heterotopic pain stimulation study in humans. *Brain Res* 2002 ; 10 : 31-40.
52. **BERMAN LH, HARTWELL GR.** Diagnosis. In : *Cohen S, Hargreaves KM, eds. Pathways of the pulp. 9th ed. St Louis: Mosby-Elsevier ; 2006.p.2-39.*
53. **GRUSHKA M, SESSLE BJ.** Application of the McGill pain questionnaire to the differentiation of toothache pain. *Pain* 1984 ; 19 : 49-50.
54. **BENDER IB.** Reversible and irreversible pulpitis : diagnosis and treatment. *Aus Endo J* 2000 ; 26 : 10-4.
55. **BENDER IB, SELTZER S, FREEDLAND, JB.** The relationship of systemic disease to endodontic failures and treatment procedures. *Oral Surg* 1963 ; 16 : 1102-15.
56. **TURK DC, DWORKIN RH, BURKE LB, et al.** Developing patient-reported outcome measures for pain clinical trials: IMMPACT recommendations. *Pain* 2006 ; 125 : 208-15.
57. **DIONNE RA, BARTOSHUK L, MOGIL J, WITTER J.** Individual responder analyses for pain: does one pain scale fit all ? *Trends Pharmacol Sc* 2005 ; 26 : 125-30.
58. **LEKIC D, CENIC D.** Pain and tooth pulp evoked potentials. *Clin Electroencephalogr* 1992 ; 23 : 37-46.
59. **KHAN AA, MCCREARY B, OWATZ CB, et al.** The development of a diagnostic instrument for the measurement of mechanical allodynia. *J Endod* 2007 ; 33 : 663-6.
60. **L.K. BAKLAND, J.O ANDREASEN** Dental traumatology : essential diagnosis and treatment planning. *Endodontic Topics* 2004 ; 7 : 14-34.
61. **ANDREASEN JO, ANDREASEN FM, BAKLAND LK, FLORES MT.** Emergency record for acute dental trauma, and clinical examination form for the time of injury and follow-up examination. In : *Traumatic Dental Injuries: A Manual, 2nd edn. Oxford : Blackwell Munksgaard ; 2003. p. 72-5.*
62. **GOPIKRISHNA V, TINAGUPTA K, KANDASWAMY D.** Comparison of electrical, thermal, and pulse oximetry methods for assessing pulp vitality in recently traumatized teeth. *J Endod* 2007 ; 33 : 531-5.
63. **EHRMANN EH.** Pulp testers and pulp testing with particular reference to the use of dry ice. *Aust Dent J* 1977 ; 22 : 272-9.
64. **TORABINEJAD M.** Pulp and periradicular pathosis. In : *Walton RE, Torabinejad M, eds. Principles and practice of endodontics. 3rd ed. Philadelphia: WB Saunders ; 2002. p.34-7.*
65. **LINSUWANONT P, PALAMARA J, MESSER H.** An investigation of thermal stimulation in intact teeth. *Arch Oral Biol* 2007 ; 52 : 218-27.
66. **SLUTZKY-GOLDBERG I, TESIS I, SLUTZKY H, HELING I.** Evidenced-based review of clinical studies on endodontic diagnosis. *Quintessence Int* 2009 ; 40 : 13-8.
67. **KLEIN H.** Pulp responses to electric pulp stimulator in the developing permanent anterior dentition. *J Dent Child* 1978 ; 45 : 199-202.
68. **KARIBE H, OHIDE Y, KOHNO H, SUGIYAMA H, UENO M, TAKAGI M, et al.** Study on thermal pulp testing of immature permanent teeth. *Shigaku Odontol* 1989 ; 77 : 1006-13.
69. **THE INTERNATIONAL ASSOCIATION OF DENTAL TRAUMATOLOGY.**

Guidelines for the evaluation and management of traumatic dental injuries. *Dental Traumatol* 2001 ; 17 : 1-4, 49-52, 97-102, 145-8.

70. **TORABINEJAD M, WALTON RE.** Periradicular lesions. In: *Ingle JI, Bakland LK, eds. Endodontics. 4th ed. Baltimore: Williams & Wilkins; 1994. p. 439-64.*
71. **COHENCA N, KARNI S, ROTSTEIN I.** Transient apical breakdown following tooth luxation. *Dent Traumatol* 2003 ; 19 : 289-91.
72. **PRIEBE WA, LAZANSKY JP, WUEHRMANN AH.** The value of the roentgenographic film in the differential diagnosis of periapical lesions. *Oral Surg Oral Med Oral Pathol* 1954 ; 7 : 979-83.
73. **BOHAY RN.** The sensitivity, specificity, and reliability of radiographic periapical diagnosis of posterior teeth. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2000 ; 89 : 639-42.
74. **BENDER IB, SELTZER S.** Roentgenographic and direct observation of experimental lesions in bone: II. 1961. *J Endod* 2003 ; 29 : 707-12.
75. **ANDREASEN FM, ANDREASEN JO.** Examination and diagnosis of dental injuries. In : *Andreasen JO, Andreasen FM, eds. Textbook and Color Atlas of Traumatic Injuries to the Teeth, 3rd edn. Copenhagen : Munksgaard ; 1993 p. 196-215.*
76. **BRYNOLF I.** Roentgenologic periapical diagnosis. IV. When is one roentgenogram not sufficient. *Swed Dent J* 1970 ; 63 : 415-23.
77. **VELVART P, HECKER H, TILLINGER G.** Detection of the apical lesion and the mandibular canal in conventional radiography and computed tomography. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2001 ; 92 : 682-8.
78. **TRONSTAD L.** Root resorption-etiology, terminology, and clinical manifestations. *Endod Dent Traumatol* 1988 ; 4 : 241.
79. **NAIR MK, NAIR UP.** Digital and advanced imaging in endodontics: a review. *J Endod* 2007 ; 33 : 1-6.
80. **MIKROGEORGIS G, LYROUDIA K, MOLYVDAS I, NIKOLAIDIS N, PITAS I.** Digital radiograph registration and subtraction: a useful tool for the evaluation of the progress of chronic apical periodontitis. *J Endod* 2004 ; 30 : 513-7.
81. **COTTI E, CAMPISI G, GARAU V, PUDDU G.** A new technique for the study of periapical bone lesions: ultrasound real-time imaging. *Int Endod J* 2002 ; 35 : 142-52.
82. **COTTON TP, GEISLER TM, HOLDEN DT, SCHWARTZ SA, SCHINDLER WG.** Endodontic applications of cone-beam volumetric tomography. *J Endod* 2007 ; 33 : 1121-32.
83. **LOFTHAG-HANSEN S, HUUMONEN S, GRÖNDAHL K, GRÖNDAHL HG.** Limited conebeam CT and intraoral radiography for the diagnosis of periapical pathology. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2007 ; 103 : 114-9.
84. **SJOGREN U, FIGDOR D, PERSSON S, SUNDDQVIST G.** Influence of infection at the time of root filling on the outcome of endodontic treatment of teeth with apical periodontitis. *Int Endod J* 1997 ; 30 : 297-306.
85. **TROPE M, BLANCO L, CHIVIAN N, SIGURDSSON A.** The role of endodontics after dental traumatic injuries. In : *Cohen S, Hargreaves KM, eds. Pathways of the pulp. 9th ed. St Louis : Mosby-Elsevier ; 2006 : 610-49.*
86. **FANIBULLDA KB.** A laboratory study to investigate the differentiation of pulp vitality in human teeth by temperature measurement. *I Dent* 1985 ; J3 (1) : 295-303.
87. **JAFARZADEH H, UDOYE CI, KINOSHITA JI.** The Application of tooth temperature measurement in endodontic diagnosis: a review. *J Endod* 2008 ; 34 : 1435-40.
88. **KIM S, SCHUESSLER G, CHIEN S.** Measurement of blood flow in the dental pulp of dogs with the 133 xenon washout method. *Arch Oral Biol* 1983 ; 28 : 501-5.
89. **TONDER KH, AUKLAND K.** Blood flow in the dental pulp in dogs measured by local H2 gas desaturation technique. *Arch Oral Biol* 1975 ; 20 : 73-9.
90. **POLAT S, ER K, AKPINAR KE, POLAT NT.** The sources of laser Doppler blood-flow signals recorded from vital and root canal treated teeth. *Arch Oral Biol* 2004 ; 49 : 53-7.

• A review of questions... •

91. RADHAKRISHNAN S, MUNSHI AK, HEGDE AM. Pulse oximetry : a diagnostic instrument in pulpal vitality testing. *J Clin Pediatr Dent* 2002 ; 26 : 141-5.
92. MIWA Z, IKAWA M, IIJIMA H, SAITO M, TAKAGI Y. Pulpal blood flow in vital and nonvital young permanent teeth measured by transmitted-light photoplethysmography : a pilot study. *Pediatr Dent* 2002 ; 24 : 594-8.
93. SIGURDSSON A. Clinical manifestations and diagnosis. In : Ørstavik D, Pitt-Ford TR, eds. *Essential endodontology*. 2nd ed. Oxford: Blackwell Munksgaard ; 2008. p. 235-61.
94. HOLLOWAY GA, WATKINS DW. Laser Doppler measurement of cutaneous blood flow. *J Invest Dermatol* 1977 ; 69 : 306-12.
95. ODOR TM, FORD TR, MCDONALD F. Effect of probe design and bandwidth on laser Doppler readings from vital and root filled teeth. *Med Eng Phys* 1996 ; 18 : 359-64.
96. YANPISSET K, VONGSAVAN N, SIGURDSSON A, TROPE M. Efficacy of laser Doppler flowmetry for the diagnosis of revascularization of reimplanted immature dog teeth. *Dental Traumatol* 2001 ; 17 : 63-70.
97. GAZELIUS B, OLGART L, EDWALL B. Restored vitality in luxated teeth assessed by laser Doppler flow meter. *Endod Dent Traumatol* 1988 ; 4 : 265-8.
98. MESAROS SV, TROPE M. Revascularization of traumatized teeth assessed by laser Doppler flowmetry: a case report. *Endod Dent Traumatol* 1997 ; 13 : 24-30.
99. RITTER ALS, RITTER AV, MURRAH V, SIGURDSSON A, TROPE M. Pulp revascularization of replanted immature dog teeth after treatment with minocycline and doxycycline assessed by laser Doppler flowmetry, radiography, and histology. *Dent Traumatol* 2004 ; 20 : 75-84.
100. EMSHOFF R, EMSHOFF I, MOSEHEN I, STROBL H. Laser Doppler flow measurements of pulpal blood flow and severity of dental injury. *Int Endod J* 2004 ; 37 : 463-7.
101. BOWES WA. Pulse oximetry : a review of the theory, accuracy and clinical applications. *Obstet Gynecol* 1989 ; 74 : 541-6.
102. SCHNETTLER JM, WALLACE JA. Pulse Oximetry as a diagnostic tool of pulp vitality. *J Endod* 1991 ; 17 (10) : 488-90.
103. SELTZER S. Alteration of human pain thresholds by nutritional manipulation and L-tryptophan supplementation. *Pain* 1982 ; 13 : 385-93.
104. GOPIKRISHNA V, PRADEEP G, VENKATESHBABU N. Assessment of pulp vitality: a review. *Int J Paediatr Dent* 2009 ; 19 : 3-15.
105. EVANS D, REID J, STRANG R, STIRRUPS D. A comparison of laser Doppler flowmetry with other methods of assessing the vitality of traumatized anterior teeth. *Endod Dent Traumatol* 1999 ; 15 : 284-90.
106. SIGURDSSON A. Pulpal diagnosis. *Endod Top* 2003 ; 5 : 12-25.
107. MORSE DR, PATNIK JW, SCHACTERLE GR. Electrophoretic differentiation of radicular cysts and granulomas. *Oral Surg Oral Med Oral Pathol* 1973 ; 35 : 249-64.
108. SKAUG N. Soluble proteins in fluid from non-keratinizing jaw cysts in man. *Int J Oral Surg* 1977 ; 6 : 107-21.
109. BELMAR M, PABST C, MARTINEZ B, et al. Gelatinolytic activity in gingival crevicular fluid from teeth with periapical lesions. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2008 ; 106 : 801-6.
110. EPELMAN I, MURRAY PE, GARCIA-GODOY F, KUTTLER S, NAMEROW KN. A practitioner survey of opinions toward regenerative endodontics. *J Endod* 2009 ; 35 : 1204-10.
111. KARDELIS AC, MEINBERG TA, SULTE HR, GOUND TG, MARX DB, REINHARDT RA. Effect of narcotic pain reliever on pulp tests in women. *J Endod* 2002 ; 7 : 537-39.
112. MESSER HH. Permanent restorations and the dental pulp. In : Hargreaves KM, Goodis HE, eds. *Seltzer and Bender's dental pulp*. Chicago: Quintessence Book ; 2002. p. 345-69.
113. BAKLAND LK. Endodontic considerations in dental trauma. In : Ingle JI, Bakland LK, eds. *Endodontics*. 5th ed. Hamilton, Ontario, Canada: BC Decker Inc ; 2002. p. 795-843.