CURRENT TRENDS TO MEASURE IMPLANT STABILITY

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INTRODUCTION
Osseointegration is defined as a direct bone anchorage to an implant body which can provide a foundation to support prosthesis.[1,2] Implant stability is a requisite characteristic of osseointegration. Without it, long-term success cannot be achieved. Continuous monitoring in a quantitative and objective manner is important to determine the status of implant stability. Osseointegration is also a measure of implant stability which can occur in two stages: Primary and secondary. Primary stability mostly occurs from mechanical attachment with cortical bone. Secondary stability offers biological stability through bone regeneration and remodeling.[3,4,5] Primary stability is affected by bone quality and quantity, surgical technique and implant geometry (length, diameter, surface characteristics). Secondary stability is affected by primary stability.[6]

Objective measurement of implant stability is a valuable tool for achieving consistently good results that are influenced by.[7] Good decisions about when to load
When a surgeon makes a decision about early loading, objective measurement of implant stability can be valuable. A specified degree of implant stability can serve as an inclusion criterion for immediate loading.
Advantageous protocol choice on a patient-to-patient basis
With objective measurement of implant stability, surgeons can make well-informed decisions about protocol choices on a case-by-case basis. In other words, when low implant stability measurements indicate that immediate loading will jeopardize treatment outcome, a two-step protocol can be applied. In cases where high implant stability measurements are recorded, the implant could be immediately loaded.

Situations in which it is best to unload
Objective measurement of implant stability also supports making the right decisions about unloading. Sennerby and Meredith point out that when replacing an immediately loaded temporary prosthesis with a permanent prosthesis, “low (secondary) values may be indicative of overload and ongoing failure.” To avoid failure, they suggest that surgeons should consider unloading, perhaps placing additional implants, and wait until stability values increase before loading the permanent prosthesis.

SUPPORTS GOOD COMMUNICATION AND INCREASED TRUST
Implant stability measurements can also help improve communication between surgeons and patients. When a surgeon refers to measurable values rather than subjective judgments as the basis for decision-making, it is easier to explain the treatment choices. The surgeons are also likely to appear more professional to colleagues alike and imbibe patient confidence.

PROVIDES BETTER CASE DOCUMENTATION
Objective implant stability measurements can be used to document the clinical outcome of implant treatments, which can be useful at a later stage if a problem should arise.

This review focuses on various methods to evaluate implant stability.

There are different methods to assess implant stability. They can be grouped as invasive/destructive methods and noninvasive/nondestructive methods.

Invasive/destructive methods
Following methods were included:
• Histologic/histomorphologic analysis
• Tensile test
• Push-out/pull-out test and
• Removal torque analysis.

Histomorphometric analysis
This is obtained by calculating the peri-implant bone quantity and bone-implant contact (BIC) from a dyed specimen of the implant and peri-implant bone. Accurate measurement is an advantage, but due to the invasive and destructive procedure, it is not appropriate for long-term studies. It is used in the nonclinical studies and experiments. It is assessed at pre-, intra-, and post-surgical time points.[8]

Tensile test
Tensile test was earlier measured by detaching the implant plate from the supporting bone. It was later modified by Bränemark by applying the lateral load to the implant fixture. However, they also addressed the difficulties of translating the test results to any area independent mechanical properties.[9]

Push-out/pull-out test
Push-out/pull-out test investigates the healing capabilities at the bone implant interface.[10] It measures interfacial shear strength by applying load parallel to the implant-bone interface. In the typical push-out or pull-out test [Figure 1], a cylinder-type implant is placed transcortically or intramedullarly in bone structures and then removed by applying a force parallel to the interface. The maximum load capability (or failure load) is defined as the maximum force on the force displacement plot, and the interfacial stiffness is visualized as the slope of a tangent approximately at the linear region of the force displacement curve before breakpoint. It is assessed during the healing period. However, the push-out and pull-out tests are only applicable for nonthreaded cylinder type implants, whereas most of clinically available fixtures are of threaded design, and their interfacial failures are solely dependent on shear stress without any consideration for either tensile or compressive stresses (Brunski et al. 2000, Chang et al. 2010). It is also technique sensitive.

Removal torque analysis
Removal torque analysis implant is considered stable if the reverse or unscrewing torque was >20 Ncm. However, the disadvantage is that at the time of abutment connection implant surface in the process of osseointegration may fracture under the applied torque stress.[11,12]

Reverse torque assessment; pull-out and push-out techniques are generally used only in preclinical applications and may be of value as research techniques. The clinical usage of destructive tests is limited due to ethical concerns associated with invasive nature of these methodologies.

Noninvasive/nondestructive methods for assessing implant stability
These include the following:
• The surgeon's perception
• Radiographical analysis/imaging techniques
• Cutting torque resistance (for primary stability)
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- Insertion torque measurement
- Reverse torque
- Seating torque test
- Modal analysis and Implatest
- Percussion test
- Pulsed oscillation waveform (POWF)
- Periotest
- Resonance frequency analysis (RFA): Electronic technology
- Magnetic technology.

The surgeon’s perception
One method of trying to evaluate primary stability is quite simply the perception of the surgeon. This is often based on the cutting resistance and seating torque of the implant during insertion. A perception of “good” stability may be heightened by the sensation of an abrupt stop when the implant is seated. An experienced surgeon’s perception is, of course, invaluable and should under no circumstances be discounted. One’s personal perception is difficult to communicate to others. However, most importantly, this type of measurement can only be made when the implant is inserted, it cannot be used later, for example, before loading the implant.

Imaging techniques
Imaging techniques are widely used to assess both quantity and quality of the jawbone. Following the surgery, imaging methods are used to assess the health of the implant, evaluating the bone quantity and quality changes, and estimating the crestal bone loss, which is a consequence of the osseointegration process. Numerous limitations exist with the use of a conventional radiograph alone in making an accurate, independent assessment of implant stability. Conventional periapical or panoramic views do not provide information on a facial bone level, and bone loss at this level precedes mesiodistal bone loss. At last, neither bone quality nor density can be quantified with this method. Even changes in the bone mineral cannot be radiographically detected until 40% of demineralization had occurred. Computer-assisted measurement of crestal bone level change may prove to be the most accurate radiographical information. However, this method is not convenient to use in clinical practice.

Cutting torque resistance analysis
This was developed by Johansson and Strid. It was later improved by Friberg et al. The amount of unit volume of bone removed by current fed electric motor and is measured by controlling the hand pressure during drilling at low speed. It determines areas of low density bone and quantifies bone hardness during implant osteotomy at the time of implant placement. Clinical studies showed that the highest frequency of implant failures was seen in jaws with advanced resorption and poor bone quality, often seen in the maxilla. Therefore, cutting resistance value may provide useful information in determining an optimal healing period in a given arch location with a certain bone quality.

The major limitation of cutting torque resistance analysis (CRA) is that it does not give any information on bone quality until the osteotomy site is prepared. CRA also cannot identify the lower “critical” limit of cutting torque value (i.e., the value at which the implant would be at risk).

Insertion torque measurement
Insertion torque values have been used to measure the bone quality in various parts of the jaw during implant placement. Insertion torque alone may be used as an independent stability measurement, but it may also act as a variable, affecting implant stability. In a different light, insertion torque is a mechanical parameter generally affected by a surgical procedure, implant design and bone quality at the implant site. However, it cannot assess the secondary stability by new bone formation and remodel around the implant. Hence, it cannot collect longitudinal data to assess implant stability change after placement. Furthermore, an increase in insertion torque may signify an increase in primary stability, but maximum insertion torque is produced by the pressure of implant neck on the dense cortical bone of the alveolus. Furthermore, it has been reported that if maximum insertion torque does not signify increased general bone density, it may indicate the insertion torque itself during tapping.

Reverse torque test
Reverse torque test was proposed by Roberts et al. and developed by Johansson and Alberktsson. It is used to assess the secondary stability of the implant. Implants that rotate when reverse torque is applied indicate that BIC could be destroyed. Further, it cannot quantify the degree of osseointegration as threshold limits vary among patients, implant material, bone quality and quantity. The studies showed, the stress of the applied torque may in itself be responsible for the failure. It also does not measure lateral stability that is a useful indicator for successful treatment outcome.

Seating torque test
Like insertion torque, the final seating torque gives some information about the primary stability of the implant when the implant reaches its final apico-occlusal position. It is done after implant placement.

Modal analysis
Modal analysis also termed as vibration analysis, measures
the natural frequency or displacement signal of a system in resonance, which is initiated by external steady-state waves or a transient impulse force. It can be performed in two models: Theoretical and experimental.

The theoretical modal analysis includes finite element analysis. It investigates vibrational characteristics of objects. It is done to calculate stress and strain in various anticipated bone levels. It is used in clinical studies and experiments. The experimental modal analysis is a dynamic analysis. It measures natural characteristic frequency, mode and attenuation via vibration testing. It is used in nonclinical studies in vitro approach. It provides reliable measurement.[25]

**Percussion test**

A percussion test is one of the simplest methods that can be used to estimate the level of osseointegration. This test is based upon vibrational-acoustic science and impact response theory. The clinical judgment on osseointegration is based on the sound heard upon percussion with a metallic instrument. A clearly ringing “crystal” sound indicates successful osseointegration, whereas a “dull” sound may indicate no osseointegration. However, this method heavily relies on the clinician’s experience level and subjective belief. Therefore, it cannot be used experimentally as a standardized testing method.[13,23]

**Pulsed oscillation waveform**

Kaneko[26] described the use of a POWF to analyze the mechanical vibrational characteristics of the implant-bone interface using forced excitation of a steady-state wave. POWF is based on estimation of frequency and amplitude of the vibration of the implant induced by a small pulsed force. This system consists of an acoustoelectric driver (AED), acoustoelectric receiver (AER), pulse generator and oscilloscope. Both the AED and AER consist of a piezoelectric element and a puncture needle. A multifrequency pulsed force of about 1 kHz is applied to an implant by lightly touching it with two fine needles connected with piezoelectric elements. Resonance and vibration generated from the bone-implant interface of an excited implant are picked up and displayed on an oscilloscope screen. It is used for in vitro and experimental studies. An in vitro study showed that the sensitivity of the POWF test depended on load directions and position.[7]

**Periotest**

Quantifies the mobility of an implant by measuring the reaction of the peri-implant tissues to a defined impact load. The Periotest was introduced by Schulte to perform measurements of the damping characteristics of the periodontal ligament, thus assessing the mobility of natural tooth.[27,28] Periotest® [Figures 1 and 2] uses an electro-magnetically driven and electronically controlled tapping metallic rod in a handpiece. Periotest value range from −8 (low mobility) to +50 (high mobility). It can measure the bone density at the time of implant placement and postsurgical placement of the implant. Response to a striking or “barking” is measured by a small accelerometer incorporated into the head. The reliability of this method is questionable because of poor sensitivity, susceptibility to many variables.[29]

**Resonance frequency analysis**

It was suggested by Meredith in 1998.[30] It is a noninvasive diagnostic method that measures implant stability and bone density at various time points using vibration and a principle of structural analysis. RFA [Figure 3] utilizes a small L-shaped transducer that is tightened to the implant or abutment by a screw. The transducer comprises of two ceramic elements, one
of which is vibrated by a sinusoidal signal (5–15 kHz) while the other serves as a receptor. The transducer is screwed directly to the implant body and shakes the implant at a constant input and amplitude, starting at a low frequency and increasing in pitch until the implant resonates. High frequency resonance indicates stronger bone-implant interface. It also provides baseline reading for future comparison and postsurgical placement of the implant. RFA has been widely used for clinically assessing osseointegration, as well as for prognostic evaluation. However, the latter aspect still has to be questioned.

The most recent version of RFA is a wireless gadget. A metal rod is attached to the implant with a screw connection. The rod has a small magnet attached to its top that is stimulated by magnetic impulses from a handheld electronic device. The rod mounted on the implant has two fundamental resonance frequencies; it vibrates in two directions, perpendicular to each other. One of the vibrations is in the direction where the implant is most stable and the other is in the direction where the implant is least stable.

Currently, two RFA machines are in clinical use: [Figure 4] Osstell® (integration diagnostics) and Implomates® (Bio TechOne).

**ELECTRONIC TECHNOLOGY RESONANCE FREQUENCY ANALYSIS (OSSTELL®)**

It was the first commercially available product for measuring implant stability. The electronic technology combines the transducer, computerized analysis and the excitation source into one machine. Implant stability quotient (ISQ) is the measurement unit (ISQ of 0 to 100) used. When used at the time of implant placement it provides baseline reading for future comparison and postsurgical placement of the implant. Currently, Osstell (Integration Diagnostic AB, Goteborg, Sweden), a commercialized product utilizing the concept of RFA, has translated the resonance frequency ranging from 3000 to 8500 Hz as the ISQ of 0–100.\[31\]

**MAGNETIC TECHNOLOGY RESONANCE FREQUENCY ANALYSIS (OSSTELL® MENTOR)**

The transducer has a magnetic peg on top and is fixed to implant or abutment [Figure 4]. On activation by magnetic resonance frequency probe the peg is activated, which vibrates and induces electric volt sampled by magnetic resonance frequency analyzer. Values are expressed as ISQ of 0 to 100. At the time of implant placement, it provides baseline reading for future comparison and postsurgical placement of the implant. However, this method is expensive and technique sensitive as it requires respective transducer and magnetic peg. It should maintain a distance of 1–3 mm, angle of 90°, and should be 3 mm above the soft tissue otherwise the measured value will be affected. Valderrama et al. reported in a study experimenting Osstell and Osstell Mentor that the two devices had high significant correlation.\[32,33\]
Newer methods under research and development

**Implant test conventional impulse testing**

Conventional impulse testing of an implant requires fastening an accelerometer with associated wires and connectors to the implant, striking it with a calibrated hammer, and then recording and interpreting the data. The objective of testing implants with electrical impulse methods is to characterize, analyze and monitor their signatures.

Implomates was developed by Huang. This device utilizes impact force from a transducer to excite the resonance of implant. The received signal is transferred to computer for frequency spectrum analysis (2–20 kHz). Higher frequency and sharp peak indicates stable implant while wider frequency and low peak indicates implant failure. At the time of implant placement provides baseline reading for future comparison and the most surgical placement of implant.

**Electro-mechanical impedance method**

This test[35] utilizes the electro-mechanical impedance of piezoelectric materials (work as both sensors and actuators) which is directly related to the mechanical impedance of the host structure. Piezoelectric zirconate titanate (PZT) is coupled to the monitored structure. After applying a voltage in 1 V in the kHz range, the PZT start to vibrate and any change of structural characteristics such as stiffness, damping, mass distribution, would influence the reading electrical admittance of PZT as read by impedance analyzer.

**Micro motion detecting device**

A customized loading device, consisting of a digital micrometer (Mitutoyo Absolute Digimatic, Mitutoyo America Corporation, Aurora, IL, USA) and a digital force gauge (Chatillon E-DFE-025, Chatillon Force Measurement Systems, Largo, FL, USA) (range of 10–2500 N 0.25% resolution over range) was used to determine implant micromotion. The forces were achieved by turning a dial, which controlled the height of the force gauge. This dialled in force was applied to the abutment via a lever. The digital micrometer was placed tangent to the crown of the abutment and detected the displacement after the load application.[36]

**Highly nonlinear solitary waves method**

HNSWs (highly nonlinear solitary waves) are compactly supported lumps of energy, which are formed by a balance between nonlinear and dispersive effects in intrinsically nonlinear media, such as granular materials. They are characterized by unique physical properties, such as high acoustic energy and remarkable robustness, which make them extremely useful as information carriers in nondestructive evaluation (NDE) applications. To generate and propagate HNSW, a granular crystal to function as a combined sensor and actuator, which is composed of a chain of spherical particles in contact with each other with a piezoelectric gauge embedded in selected locations. Using the granular crystal, the surface of an orthopaedic implant with a single HNSW, and record the signals reflected from the interface between the granular crystal and the implant specimen under inspection.

Here, granular crystal actuator consisting of a one-dimensional tightly packed array of spherical particles, to generate acoustic solitary waves are assembled through direct contact with the specimen. Acoustic solitary waves into a biomedical prosthesis are injected, nondestructively evaluating the mechanical integrity of the bone-prosthesis interface, studying the properties of the waves reflected from the contact zone between the granular crystal and the implant. The granular crystal contains a piezoelectric sensor to measure the traveling solitary waves, which allows it to function also as a sensor.

Then a sequence of harsh mechanical loading on the samples is imposed to degrade the mechanical integrity at the stem-cement interfaces, using simulator that simulates aggressive, accelerated physiological loading. Implant stability is investigated via the granular crystal sensor-actuator during testing. Results showed that the reflected waves respond sensitively to the degree of implant fixation. In particular, the granular crystal sensor-actuator successfully detects implant loosening at the stem-cement interface following violent cyclic loading. This technique[37] suggests that the granular crystal sensor and actuator has the potential to detect metal-cement defects in a nondestructive manner for orthopedic applications.

**SUMMARY AND CONCLUSION**

Evidence from the presented literature indicates that the advanced tests and equipment may play a greater role in the evaluation of implant stability compared to conventional methods. The ability to monitor osseointegration and the life expectancy of an implant is a valuable diagnostic and clinical tool that has far-reaching consequences on implant dentistry, RFA has attracted considerable scientific interest in recent years; it can also be used to evaluate the effect of early and delayed loading, assess stability over a period of time and early diagnosis of implant
failure. However, information should be established from many different diagnostic aids to assure long-term implant stability.

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