Embracing new technology, with caution

SADJ February 2023, Vol. 78 No.1 p49-51
LM Sykes¹, V Bookhan ²

ABSTRACT
Dental manufacturers frequently present clinicians with new “cutting edge” materials, devices or technology. These usually come with great promise for bettering the status quo in their practices, and of putting them ahead of the colleagues in the market place. However before succumbing to the advertorial hype, and abandoning their old practices, materials or equipment, practitioners need to evaluate the new offering against the “gold standard” if one exists. This entails comparing it to “the benchmark” practice / product that is routinely used under reasonable conditions, and answering a number of clinically and scientifically pertinent questions. If they are then confident it has been through extensive trials, the results have been analysed with appropriate tests by independent investigators, and the reporting thereof is accurate, reliable, repeatable, sensitive, specific and clinically applicable, they may consider making practice changes. While it is admirable for clinicians to be open minded and willing to embrace and adapt to modern technology, this should only be done if the change has been proven superior to reliable routine practices. It is incumbent on all practitioners to keep abreast of current trends through the many platforms available. They should also strive towards being life-long learners who are curious, open minded, flexible, willing to learn new skills, and open to adapting their work to embrace advances. This will hopefully lead to practitioners having more fruitful careers, and equip them to provide the best possible service and care to their patients.

INTRODUCTION
How many ways are there to peel an orange and what does this topic have to do with dentistry? The answer to both those questions is “Probably a lot more than you think”, as you will discover in this paper where the underlying fruit. The conventional hand peeling method has been carried out for centuries, and has remained in use due to its simplicity and reliance on basic skills and equipment (in this case hands). It does however require time, effort, a certain amount of manual dexterity, and follows steps that can vary with each person. It can be messy and may require the peeler to go back over sections to remove small residual remnants of rind or pith. The outcomes are generally satisfactory. However, the procedure also creates debris which differs in amount and size of rind pieces removed, seldom creates a smooth surface finish, and may not produce the most aesthetically pleasing peeled orange. The end result is largely dependant on the type and size of the orange, its condition at the beginning of the procedure and the operator’s style and dexterity. It can be a laborious and time consuming process, leaving one to wonder if the effort justifies the outcome (Figure 1).

With time and improved technology, new equipment was introduced to the culinary market. Sharp knives were marketed as multi-purpose gadgets that claimed to save time, be easy to use, be clean and comfortable to work with, remove peels and pith efficiently, and generate less debris. They gained widespread use not only for peeling oranges, but also for a myriad of other domestic purposes. All knives had a basic design of handle and sharp cutting blade and were initially cheaply to purchase. The expense seemed justified as the procedure was relatively easy and effortless in comparison to hand peeling, didn’t take much skill or practice to master, allowed the entire peel and pith to be cut off in a few strips that were easy to discard, and generally saved a lot of time (Figure 2). As their popularity grew so did technology and soon the markets were flooded with knives. The handles and blades came in different materials, colours, sizes and shapes, and were often “custom designed” for specific purposes. Knives looked set to become the “best-practice” orange peelers on the market and many people purchased them. Sadly, with time and use it became evident that the cutting process had certain drawbacks and some unwanted side effects. There was a lot of fluid spillage, many operators sustained finger injuries, they often cut too deeply into the fruit resulting in loss of the valuable fruit substance, and sometimes even damaged the underlying pulp. In addition, many of the early blades corroded from the acidic juices, and both the knife users and the fruit recipients began to complain and to even look for safer options that would also result in less damage.

Author contributions:
1. LM Sykes BSc, BDS, MDent, FCDS(SA), IRENSA, Dip Forensic Path, Dip ESMEA, Head of Department of Prosthodontics, University of Pretoria. https://orcid.org/0000-0002-2002-6238
2. Vinesh Bookhan BDS, MDent, Department of Odontology, University of Pretoria. https://orcid.org/0000-0002-4235-3897

Corresponding Author:
Name: L Sykes
Email: Leanne.sykes@up.ac.za

Authors contribution:
1. Leanne M Sykes: Primary author 90%.
2. Vinesh Bookhan : Second author 10%.

*The illiterate of the 21st century will not be those who cannot read or write, but those who cannot learn, unlearn and relearn* Alvin Toffler
Some innovative thinkers devised a new gadget specifically for peeling oranges and certain other fruits. The idea was to create a utensil that was functional, efficient, corrosion resistant, durable, easy to clean, cheap to manufacture and affordable. The new synthetic gadget had some of the knives features such as a comfortable handle and a slick cutting tip, but did away with the potentially dangerous blade. It was marketed by the manufacturers who did countrywide road shows where they generally gave an introductory talk, a live demonstration, and a short hands-on training session for interested purchasers. Features included a hooked tip to score the fruit a predetermined depth, and to cut down along planned and equally spaced vertical lines, this ensured that all pieces of rind were similarly sized and evenly thick. The curved body portion hugged the contours of the fruit as it gently separated the peel away without causing any damage to the underlying fruit. It resulted in a neat smooth finish, minimal debris, and had an added bonus of vertical grooves along which the orange could be opened (Figure 3). Introductory packages often came at good prices with “additional extras” included to sweeten the deal. These gadgets did indeed serve their intended purpose and at a much reduced price. Results were good, predictable, and the damage and debris much less. However, they had their limitations being designed for only a few specific applications, and it was incumbent of the operator to know when and where to use it.

RELEVANCE TO DENTISTRY

The above scenarios may seem far removed from dentistry, and the reader will be forgiven for asking “What does this topic have to do with me and my practice? The answer to both those questions is “Probably a lot more than you think”.

In a similar vein to the orange peeling analogy, dental manufacturers frequently present clinicians with new “cutting edge” materials, devices or technology. These usually come with great promise for bettering the status quo in their practices, and of putting them ahead of their colleagues in the market place. However before succumbing to the advertorial hype practitioners needs to take time and make an effort to do some basic investigating themselves. They should evaluate the new offering against the “gold standard” if one exists. This entails comparing it to “the benchmark” practice / product that is routinely used under reasonable conditions. It also requires them to “precisely define the question of interest (clinical question), look for relevant information about it from databases, study the research methodology that was used during development and trial periods, including the statistical analysis, critically evaluating the quality of the studies and understand their implications in terms of use and patient care”.

Thus when clinicians are presented with any new product they need to critically evaluate the associated literature presented to them by the manufacturers or sales representatives, and ask a number of pertinent questions. This includes inquiring: If the product has been tested in a laboratory? Were tests standardised and carried out according to approved protocols? What were the results of the testing? Is it biologically safe for use in patients? Is it safe for use by clinicians? Were the trials company sponsored as this can lead to an element or researcher bias? Was any conflict of interest declared in the research? Have the results been validated by other independent researchers? Have the results been published in accredited peer reviewed journals and not only in company catalogues and prints? Has the product been subject to human trials? Are there long-term follow up studies? Were there any adverse events? If so,
have they been reported? What is the cost of the new innovation? Does it have a shelf life and/or how often will it need to be replaced? Does it require training to use? If so, who will provide the training, the company, trained clinicians, or an academic institution? Is there any maintenance plan if equipment is involved and is service easily available after purchase? Only if all of these queries can be satisfactorily answered should the clinician consider replacing the old with the new.

Thereafter, if a practitioner decides to invest in the new product or to make any substantial changes in their routine work, they should proceed with caution. If they see that it is not performing as ideally as promised or anticipated they have a “duty to care” for their patients and to assess the situation in an unbiased manner. This may involve evaluation of the intervention as well as a degree of self-reflection to ensure that the shortcomings are not due to their own inadequacy or lack of training and skills. Thereafter they may need to report their observations to the manufacturers as well as to alert colleagues.

In medical research, Good Clinical Practice (GCP) guidelines have been established to protect patients’ rights, and to ensure their safety throughout any clinical trial period, including scheduled follow up evaluation. While “mostly directed towards investigators, pharmaceutical and technological manufacturers, research sponsors, trial participants, research ethics committees, and medicines regulatory authorities”, it is also incumbent on practitioners to be part of the process and to monitor newly implemented interventions. They have a moral obligation to look back on their findings if any adverse events or problems are noted. This feedback may take place in informal discussions, in small working groups, via correspondence with manufacturers and regulatory bodies, or through publication in dental newsletters and journals. They should also stop using the new regime immediately regardless of how much they have invested in the initial outlay. It would be unethical to continue to use up stock or keep working with the equipment merely to justify its expense. Practitioners may also be afraid to make their observations public in case the manufacturers accuse them of slander. This fear may be allayed if they publish their findings as observations, and ask colleagues to report if they have had any similar experiences. This will alert others to be more vigilant if they are using the new intervention, as the saying goes “You see only what you look for, you recognise only what you know”. If the adverse events are too frequent or serious in nature, then the new intervention should be rejected immediately. Furthermore, in the interest of beneficence, and good communication, the dentist may need to alert the patients of potential problems, and offer assistance if they develop complications related to this treatment.

CONCLUSION

Over the years many new products have come onto the dental market promising features such as superior strength, better bonding, reduced tooth sensitivity, enamel remineralisation, multi-purpose uses, easy manipulation, good taste, long shelf life, superior aesthetics, and a range of other desirable features. Some lived up to their promises, others were replaced by updated versions or different products, many ended up in the back of store rooms, or were completely discarded. The latter generally disappeared from sight and use, except perhaps for mention in materials textbooks, (which should be read with the awareness that they become rapidly dated). This paper highlights the need for clinicians to keep abreast of current trends and innovations in dentistry, to read trustworthy scientific literature, and to ensure that before they embark on any new treatment modality they are confident it has been through extensive trials, the results have been analysed with appropriate tests by independent investigators, and the reporting thereof is accurate, reliable, repeatable, sensitive, specific and clinically applicable. While it is admirable for clinicians to be open minded and willing to embrace and adapt to modern technology, this should only be done if the change has been proven superior to reliable routine practices. It is incumbent on all practitioners to spend time reading contemporary literature, attending congresses, participating in study groups, and contributing information based on their own experiences to colleagues in the field. Life-long learning and communication will hopefully lead to practitioners having more fruitful careers, and equip them to provide the best possible service and care to their patients.

REFERENCES