

Characterisation and Physico-Chemical Evaluation of a Novel Glass Ionomer Nanozirconia- Silica-Hydroxyapatite Hybrid Material



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INDUTROCTION

Glass ionomer cement (GIC) is among the popularly used materials prosthodontics for core build- up and cementation of prostheses. However, GICs suffer from relatively inferior mechanical properties and sensitivity to initial desiccation. A novel GIC-nano Zirconia-Silica- Hydroxyapatite (GIC-nano ZrO₂-SiO₂-HA) has been developed which has better hardness and aesthetics (1-4).

OBJECTIVES

To characterise and evaluate the colour stability and fluoride release of a new GICnano ZrO₂-SiO₂-HA hybrid material.

METHODOLOGY



- The nano ZrO₂-SiO₂-HA was synthesised using modified sol-gel technique (3), and the nanopowder was characterised using TEM, FTIR and XRD.
- The samples (*n*=10) per group were prepared by adding 5% by weight of nano ZrO₂-SiO₂-HA powder into conventional GIC (Fuji IX).
- The colour stability was measured (ΔE) with a spectrophotometer using the CIE L*a*b* system, and the fluoride (F) release was measured using an ISE meter over a one-month period.

CHARACTERISATION TEM MICROGRAPHS



- Peak at 564.78- Refers to v₄ bending mode of PO_4^{3-} indicative of HA.
- 604.06- Assigned to O Si O and Zr –O Zr bending and presence of HA.
- 875.11- Signifies HA with deficiency of Ca.
- 963.31- Certifies the structure of ZrO₂.
- Therefore, it can be inferred that a homogenous nano ZrO₂-SiO₂-HA powder was successfully synthesised using sol-ge technique(3).

XRD DIFFRACTOGRAM



- ~25° and 28°- 002 and 102 plane of HA; 011 and 111 plane of Sio₂, respectively.
- ~32°and ~34°- 011 and 002 plane of ZrO₂, respectively.
- ~40°- 212 plane of HA.
- ~59 60°-121 plane of ZrO₂.
- These findings are in agreement with the results of FTIR spectroscopy.

RESULTS COLOUR STABILITY

FLUORIDE RELEASE



Both the control and GIC nano ZrO₂-SiO₂-HA showed similar patterns of F release during the course of the study. Repeated measures ANOVA reveals a highly significant difference (p≤0.05) in mean F release between both the groups for all the time intervals except for day 2, day 4 and day 28.

CONCLUSION

- morphologic and qualitative The characterisations of the nano ZrO₂-SiO₂-HA synthesised by a modified one-pot synthesis confirmed the presence of homogenously incorporated functional groups corresponding to each element.
- All the particles were in the nano-scale range, with spherical ZrO₂ and SiO₂ particles embedded in the voids between rod-shaped HA crystallites enhancing the packing density.
- The colour change between various time intervals shows that the ΔE values for GIC $nanoZrO_2$ -SiO_2-HA were < 3.3 (i.e. acceptable) and generally were lower than those of the control group. Therefore, GIC nano Zr-Si-HA exhibited enhanced colour stability.
- It can be concluded that the addition of nano ZrO₂-SiO₂-HA to the GIC did not impede its F⁻ releasing ability. It in fact resulted in an overall higher F⁻ elusion from the GIC nanoZrO₂-SiO₂-HA when compared to the control group.

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TEM micrographs confirmed that all of the powder particles were in the nano scale range with the rod-shaped HA particles interspersed with spherical nano ZrO_2 and $SiO_2(1,3)$.



Repeated measures ANOVA shows a statistically significant difference between the day 1 to day 7 (ΔE_1) and day 14 to day 28 (ΔE_2) values for control (Fuji IX) and GIC 5% nano ZrO₂-SiO₂-HA. However, $\Delta E_2(1.09)$ and $\Delta E_3(1.32)$ for GIC 5% nano ZrO₂-SiO₂-HA were much lower compared to the control group ($p \le 0.05$).

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