

1. Introduction

The use of composite resins as restorative materials in Dentistry has been extensive. However, the toxicity has been linked to some of its components, in particular the monomers present as a result of incomplete polymerization, which may elute into the oral cavity^{1,2}. The appearance on the market of a new range of resins that calls the possibility of restoration block with thicknesses up to 4 mm requires further studies in vitro and in vivo to ensure its biocompatibility.

2. Objective

To compare the cytotoxic reaction produced by two different composite resins Bulk fill, namely Filtek™ Bulk Fill (3M™ ESPE) (FBF) and Tetric EvoCeram® Bulk Fill (Ivoclar Vivadent®) (TEC), used in dental restoration, in cultures of fibroblasts in order to infer about their biocompatibility.

3. Methods

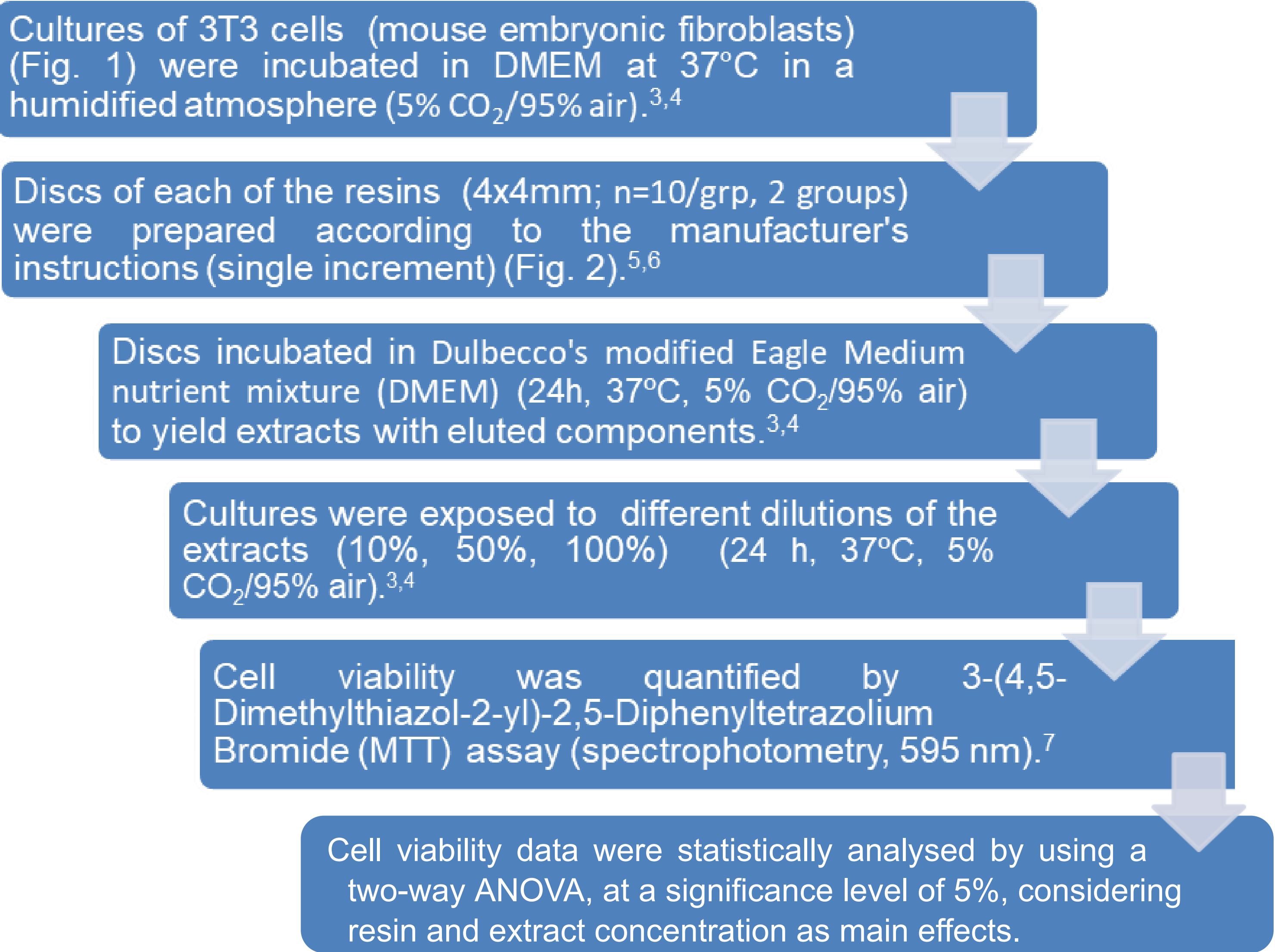


Fig. 1 — 3T3 fibroblasts observed at the microscope: 1 day after thawing (left); With 80% confluence after 3 days of incubation in DMEM - used in cytotoxicity tests (center); Morphological changes visible after 24h of contact with FBF100% (right).

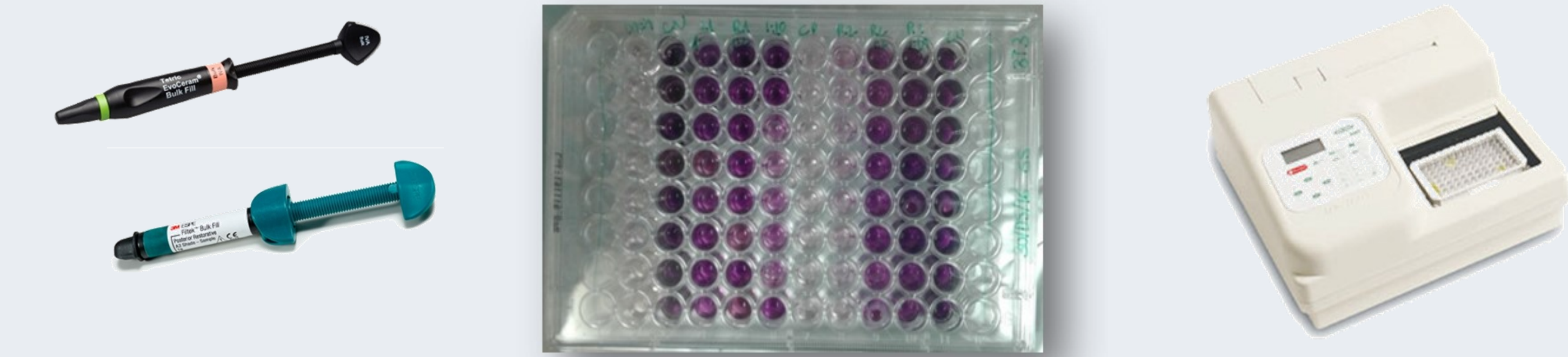


Fig. 2 — TEC and FBF resin-composites^{5,6} used in this study (left); The 96-well plate after incubation of the cells for 24 h with the resin extracts and solubilization of the formazan crystals with dimethyl sulfoxide (DMSO), immediately prior to the spectrophotometric analysis (center); Spectrophotometer microplate reader, model 680 (Bio Rad®) (center).

Bibliography

1) Geurtsen, W. (2000). Biocompatibility of Resin-Modified Filling Materials. *Critical Reviews in Oral Biology & Medicine*, 11(3), 333–355.
2) Van Landuyt, K. L., et al. (2011). How much do resin-based dental materials release? A meta-analytical approach. *Dental Materials*, 27(8), 723–747.
3) International Organization for Standardization. (2008). Evaluation of biocompatibility of medical devices used in dentistry.
4) International Organization for Standardization. (2009). Biological evaluation of medical devices. Part 5: In vitro cytotoxicity tests (ISO 10993-5: 2009)
5) 3M ESPE. (2015). Filtek™ Bulk Fill - Scientific Documentation.
6) Ivoclar Vivadent (2016). Tetric EvoCeram Bulk Fill, Scientific Documentation. Instructions for use
7) Vitral, J. C. D. A. (2008). Avaliação da Citotoxicidade de Materiais Odontológicos Através do Método de MTT e Produção de Óxido Nítrico: Descrição de uma Técnica. *Pesquisa Brasileira Em Odontopediatria E Clínica Integrada*, 8(3), 359–365.

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4. Results

Concentration of the extract	Citotoxicity (%)	
	TEC	FBF
100%	72,0	81,1
50%	49,2	39,6
10%	41,5	36,9

Table 1 - Comparative cytotoxicity between the TEC and FBF resins at different dilutions of extract.

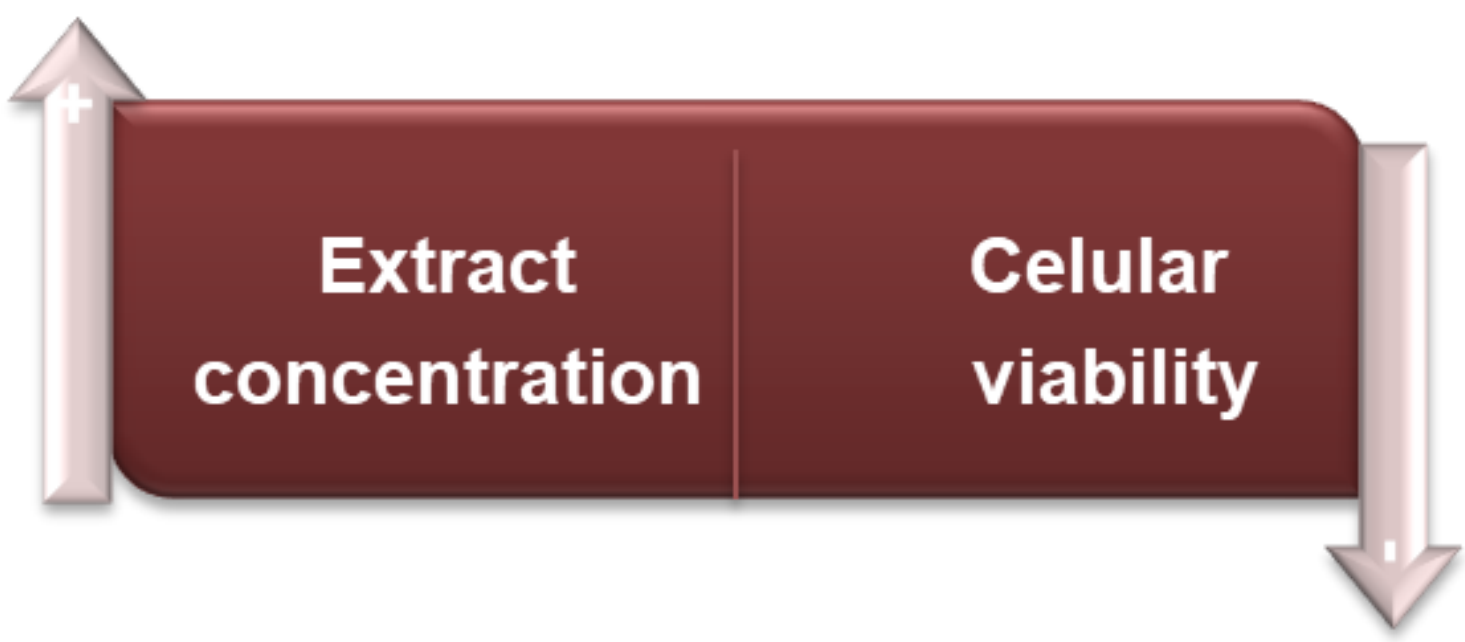
All dilutions of added resin extracts caused high cell death.

No statistically significant differences were observed between the cytotoxicity of the tested resins (p=0.154).

In both resins, there was a decrease in cell viability when increasing extract concentration (p=0.809).

There's a correlation between extract concentration and toxicity.

No interaction effect was identified between resin and extract concentration (p=0.809).



5. Conclusions

- In the first 24h, there was release of cytotoxic components in both resins.
- They were cytotoxic in all dilutions tested, although no significant differences were observed between them.
- The identification of toxic components is important to avoid their use in dentistry.