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F8 A PROSPECTIVE, RANDOMIZED, DOUBLE-BLINDED TRIAL TO STUDY THE EFFICACY OF TOPICAL TOCOTRIENOL IN THE PREVENTION OF HYPERTROPHIC SCARS

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Background: Despite widespread beliefs regarding the use of topical tocotrienol in the prevention of hypertrophic scars, there is very little evidence from well-controlled and randomized clinical trials to justify its benefits for surgical scars.

Objective: The present study was conducted to evaluate the efficacy of topical tocotrienol in preventing the development of hypertrophic scars.

Methods: A prospective, randomized, double-blinded study was performed in which the patients were randomized into either treatment group with 5% topical tocotrienol or a placebo group. The patients were required to apply the preparation to their scars twice a day for six weeks starting at two weeks after surgery. An evaluation of the scars was performed at weeks 0, 2, 6 and 16 following the onset of topical application using three methods: a clinical assessment using the Patient and Observer Scar Assessment Scale (POSAS), a photographic scar assessment by two independent assessors using a visual analogue scale, and laser Doppler imaging (LDI).

Results: There was no statistically significant difference in scar parameters between the tocotrienol group (n=43) and the placebo group (n=40) in the POSAS, photographic scar assessment or mean flux of LDI ($p>0.05$). The mean LDI flux showed decreasing trend over time, which was positively correlated with the vascularity score (correlation coefficient=0.322) of observer scar assessment scale on week 0 ($p=0.018$) and total score of the patient scar assessment scale on week 6 (correlation coefficient=0.354, $p=0.009$). No significant adverse effects were observed.

Conclusions: Twice daily application of 5% topical tocotrienol had no significant effect on the appearance and perfusion of scars over four months post-surgery. LDI has a promising role as a scar assessment tool.