

A Prospective, Randomized, Controlled Trial Comparing the Outpatient Treatment of Pediatric and Adult Partial-Thickness Burns with Suprathel or Mepilex Ag

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Indication:	Partial-thickness burns (mean TBSA 5.5±4.6%)
Study Design:	Prospective, randomized, controlled,
Treatment:	Suprathel® in comparison to Mepilex® Ag
Group sizes:	Suprathel®: n=32; Mepilex® Ag: n=30; pediatric subgroup: n=33

Pain » Pain ratings were significantly lower in Suprathel® group compared to Mepilex® Ag group during first 5 days.

Scarring » POSAS (Patient and Observer Scar Assessment Scale) was significantly reduced in burns by 56% compared to Mepilex® Ag after 1 month.

