LAVILON
Lotion Antichute

Double Blind Clinical Trial for the evaluation of efficacy, skin tolerance and cosmetic acceptability of LAVILON LOTION ANTICHUTE compared with a similar reference product.

100 subjects with androgenetic alopecia

(two groups of 70 males and 30 females; Age 30 - 50 years old).

Treatment duration: 3 months in comparison with a similar reference product

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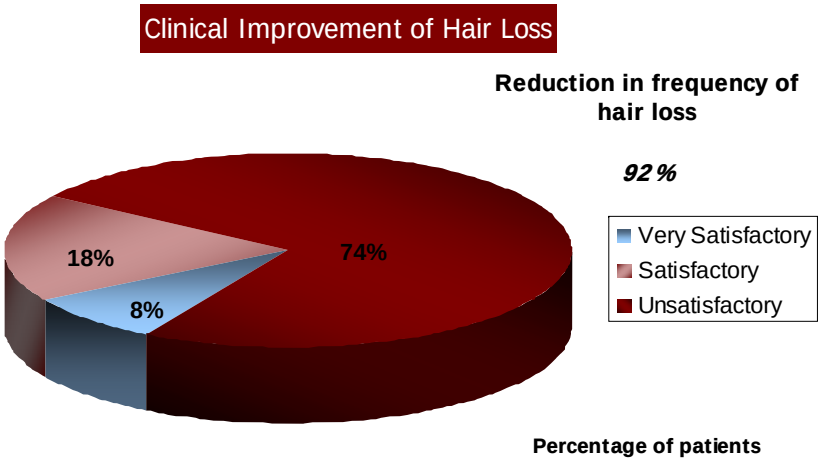

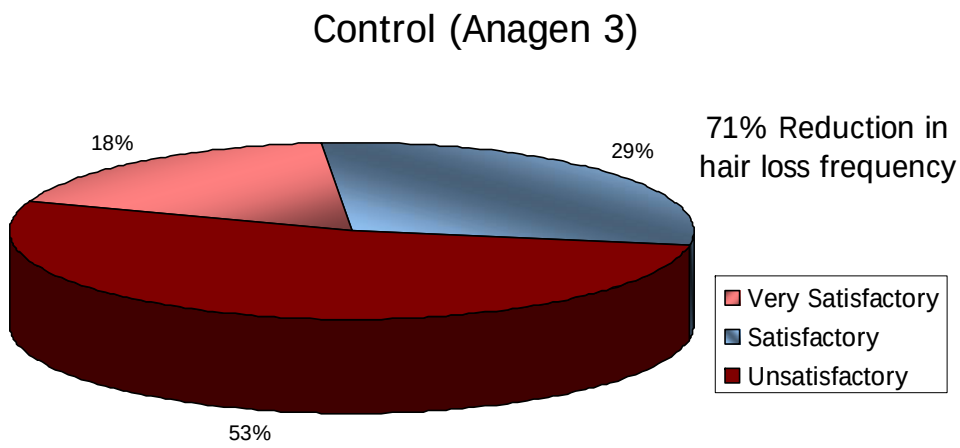

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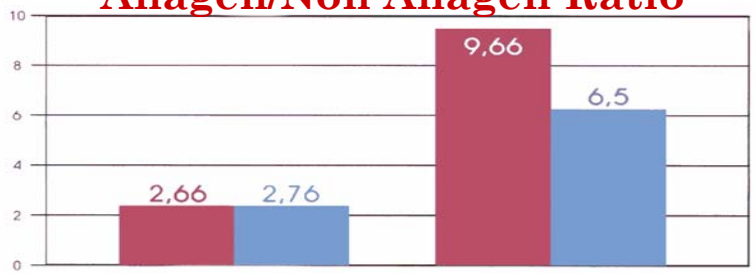
Promotion of hair in anagen phase

Anagen/Non Anagen Ratio



70%

Promotion of healthy hair in anagen phase

Trichogram

Increased diameter of shaft

	Mean shaft diameter (mm)	
LOTION ANTICHUTE	0.040	0.095 p<0.05
Control (Anagen 3)	0.040	0.065 p=n.s.

138%

Increase in hair density

Greater cosmetic acceptability

Excellent tolerability

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