

## Certificate of Analysis

(Representative Sample Certificate)

**Product Name:** Follicle Booster  
**INCI Name:** Butylene glycol, water, dextran, acetyl tetrapeptide-3, Trifolium Pratense (Clover) flower extract  
**CAS Number:** 107-88-0, 7732-18-5, 9004-54-0, 827306-88-7, 85085-25-2  
**Lot Number:** Not available (data may vary slightly with different lots or batches)  
**Expiration Date:** 36 months from production date

Property	Specification	Analysis
Appearance (Visual)	Transparent liquid	Pass
Color (Visual)	Colorless	Pass
Odor (Olfactive)	Characteristic	Pass
pH (USP <791>)	4.0-6.0	5.3
Refractive Index (USP <831>)	1.380-1.415	1.393
Identification of Active Ingredient (Acetyl Tetrapeptide-3) (HPLC Method)	Acetyl Tetrapeptide-3	Acetyl Tetrapeptide-3
Dosage of Active Ingredient (Acetyl Tetrapeptide-3) (HPLC Method)	≥240 ppm	295 ppm
Identification of Active Ingredient (Biochanin A) (HPLC Method)	Biochanin A	Biochanin A
Dosage of Active Ingredient	≥15 ppm	37 ppm

**Disclaimer:** This information relates only to the specific material designated and may not be valid for such material used in combination with any other materials or in any other process. Such information is to be the best of the company's knowledge and believed accurate and reliable as of the date indicated. However, no representation, warranty or guarantee of any kind, express or implied, is made as to its accuracy, reliability or completeness and we assume no responsibility for any loss, damage or expense, direct or consequential, arising out of use. It is the user's responsibility to satisfy himself as to the suitability & completeness of such information for his own particular use.

(Biochanin A) (HPLC Method)		
Total Plate Count (USP <61>)	<100 CFU/mL	<10 CFU/mL
Yeasts & Molds (USP <61>)	<10 CFU/mL	<10 CFU/mL
E. coli (USP <62>)	Absence	Absence
S. aureus (USP <62>)	Absence	Absence
P. aeruginosa (USP <62>)	Absence	Absence
C. albicans (USP <62>)	Absence	Absence

The above data were obtained using the test indicated and is subject to the deviation inherent in the test method. Results may vary under other test methods or conditions.

This report is not to be signed.

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