

## **ANALYTICAL LABORATORIES**

microbiology - physicochemistry - sensory





## TEST REPORT No.: L/0/08/2021/292/M/1/EN

Order No.:

Ł/0/08/2021/292

- A accredited methodology (AB 1095); reference if the law so provides (the result can be used to assess compliance in the legally regulated area).
- AE accredited methodology (AB 1095) of flexible scope reference if the law so provides / equivalent to reference (the result can be used to assess compliance in the legally regulated
- AR accredited methodology (AB 1095) equivalent to reference (the result can be used to assess compliance in the legally regulated area).
- MON methodology accredited in terms of "OiB"
- GMP+ methodology registered in the scope of GMP+ B11 protocol (feed testing)
  - A/P accredited methodology of the subcontractor
    - P non-accredited methodology of the subcontractor

Materia	al/product tested: Cosmetics							
Produc	t name: PROPAR	IN - Sham	poo Hair	Growth System		Date: 5 si	erpnia 2021	
Produce	er:	ov	vn produc	tion				
Date of	production:	no	data					
Lot nun	•	06	.2021/34	14				
Notes o	n the sample:	Pr	eserving s	system: - Phenoxyethanol - 0.823:	5% - Ethylhexylglycerin - 0.1035	%		
	collected according to: transported by: Shipping				Sample receiver:	JARS employee n	ю.: 2533	
Sample	no.: 4855/08/21 Sample evaluation	un	reservedly	y Analysis start da	te: 11-08-2021 Analy	rsis end date:	13-09-202	1
Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	N
	Preservation efficiency - challenge test	-	AE	PN-EN ISO 11930:2019-03		in Attachment		
Ł								

<sup>\*\* -</sup> expanded measurement uncertainty at the level of confidence app. 95% and the coverage factor k=2, does not take into account the sampling uncertainty, except when indicated in the remarks.

Measurement uncertainty of the test results is presented when: it is relevant to the validity or application of the test results, it affects conformity to a specification limit, or a customer's instruction so requires.

#### Remarks:

NOTE: The original test reports are issued as PDF file, signed with a qualified electronic signature. Therefore, all prints are copies, unless certified to be true to the original PDF file.

Report prepared in a single copy		The end of the Report	Original of PDF: Customer, copy of PDF to: Laboratory archive
Created on:	Authorized by:	Approved by:	
08-10-2021	JARS employee no.: 2275	Specjalista ds.Kosm	netyków Signed with a qualified electronic signature
		JARS employee no.	.: 2098

1/1 Ł/0/08/2021/292/M/1/EN

The results relate to the tested samples.

The test report includes test results of the following number of samples: 1 pc(s) and without the written approval of the laboratory shall not be reproduced except in full.

Customer may file complains within 14 days from receiving the report.

The laboratory does not store the samples after testing, unless otherwise agreed with the customer.

Place of performance of the tests (location codes): L - Lajski, M - Mysłowice, PS - in situ measurement.



## Attachment no M/4855/08/21 to Test Report

## Preservative Test according to PN-EN ISO 11930:2019-03

Sample number: 4855/08/21

		Number of		Results unit			
Microbial s	train	microorganisms in standarized suspension	Contamination level	After 7 days	After 14 days	After 28 days	
Pseudomonas ac	eruginosa	N-4.0-408 -5./1	N =4 6×405 of v/a	N <sub>x</sub> =<1,0x101 cfu/g	$N_x = <1,0x10^1 \text{ cfu/g}$	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	
ATCC 902		N=1,6x10 <sup>8</sup> cfu/ml	N₀=1,6x10⁵ cfu/g	Log R = ≥4,2	Log R = ≥4,2	Log R = ≥4,2	
Staphylococcus	s aureus	N. 4.0.408 -5./1	N -4 0-405 af-/a	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	
ATCC 6538		N=1,8x108 cfu/ml	N₀=1,8x10 <sup>5</sup> cfu/g	Log R = ≥4,3	Log R = ≥4,3	Log R = ≥4,3	
Escherichia coli		N=1,7x108 cfu/ml	No=1,7x10 <sup>5</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	
	ATCC 8739			Log R = ≥4,2	Log R = ≥4,2	Log R = ≥4,2	
Candida alb	icans	N=6,0x10 <sup>7</sup> cfu/ml	N <sub>o</sub> =6 0x10 <sup>4</sup> cfu/a	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	
ATCC 102				Log R = ≥3,8	Log R = ≥3,8	Log R = ≥3,8	
Aspergillus bra	siliensis	N 0 0 407 5 4-1	N₀=6,6x10 <sup>4</sup> cfu/g		N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	N <sub>x</sub> =<1,0x10 <sup>1</sup> cfu/g	
ATCC 164		N=6,6x10 <sup>7</sup> cfu/ml		X	Log R = ≥3,8	Log R = ≥3,8	
(A)		Bacteria	≥3	≥3 and NI	≥3 and NI		
Requirements <sup>a</sup>		Log reduction values (A)	Candida albicans	≥1	≥1 and NI	≥1 and NI	
		Log	Aspergillus brasiliensis	x	≥0 <sup>b</sup>	≥1 and NI	

a - acceptable aberrance from acceptable value degree reduction number of microorganisms is 0,5 log, cfu- colony forming unit, NI- no increase from the previous count;  $N_0$ =number of cfu in 1g of sample after contamination,  $N_x$ = number of cfu in standardized suspension,  $N_x$  – number of cfu in 1 g sample after a certain time; Log R= Reduction values in Log where R=  $N_0$ -  $N_x$ , x- not performed, NI- the number of microorganisms no increase in the count from the previous contact time, b- the number of microorganisms no increase in the count from the initial contact time

#### Demonstration of the neutralizer efficacy

Microbial strain		Res	ults	
Microbial strain	Nvf / Nvn	Nvn	Nv	Requirements
Pseudomonas aeruginosa ATCC 9027	0,70	111	113	
Staphylococcus aureus ATCC 6538	0,71	122	128	Nvf / Nvn ≥ 0,5 Nvn ≈ Nv
Escherichia coli ATCC 8739	0,77	117	123	
Candida albicans ATCC 10231	0,77	52	58	
Aspergillus brasiliensis ATCC 16404	0,70	57	61	

Nv- counts of the bacteria performer on the inoculum, Nvn – counts of the bacteria performed on the mixture of the neutralizer and diluents, Nvf - counts of the bacteria performed on the mixture of the neutralizer and the formulation

#### Comments:

## ADDITIONAL INFORMATION:

- · Preliminary tests: microbiologically pure product
- · Neutralizer: D/E broth
- Tryptic soy agar 32,5°C±2,5°C/3 days (bacteria), Sabouraud dextrose agar 32,5°C±2,5°C/3 days (Candida albicans), Potatoe Dextrose Agar 22,5°C±2,5°C/5 days (Aspergillus brasiliensis)

#### **END of Attachment**

Prepared on:	Prepared by:	Authorized by:
8.10.2021	JARS employee no 2408	JARS employee no 2275



## Attachment No. F/12435/07 to the Test report

Product name:

PROPARIN - Shampoo Har Growth System

Sample number:

12435/07/21

Test type:

Dermatological (patch test)

Date of test start:

21.07.2021

Date of test end:

28.07.2021

#### Qualitative product composition:

INCI composition (\*): Aqua, Sodium Coco-Sulfate, Sodium Cocoamphoacetate, Cocamidopropyl Betaine, Glycerin, Polysorbate 80, Sodium Chloride, Sodium Lauroyl Sarcosinate, Cetrimonium Chloride, Polyquaternium-10, Betaine, Ecklonia Cava Extract, Larix Europaea Wood Extract, Camellia Sinensis Leaf Extract, Biotinoyl Tripeptide-1, Apigenin, Glycine, Coffea Arabica Seed Extract, Ethoxydiglycol, Zinc Chloride, Oleanolic Acid, Panthenol, Pantolactone, Parfum, Hydroxypropyl Methylcellulose, Potassium Sorbate, Lactic Acid, Butylene Glycol, 1,2-Hexanediol, Sodium Benzoate,PEG-40 Hydrogenated Castor Oil, PPG-26-Buteth-26, Phenoxyethanol, Ethylhexylglycerin, Disodium EDTA.

(\*) - The Customer bears full responsibility for compliance of samples delivered for testing with the declared qualitative composition. The Laboratory does not analyse the composition in regard to the current regulatory requirements.

## Microbial purity testing

Microbiologically approved product.

The microbiological purity tests carried out in JARS Laboratory / delivered by the Customer.

#### Dermatological tests

Patch dermatological tests / contact tests are used to detect contact allergy (delayed type allergy) to small molecule compounds (haptens). Sensitization occurs through contact with the antigen that penetrates the skin. The antigen is recognized by Largenhans cells and presented to T lymphocytes. Specifically sensitized lymphocytes multiply in the regional lymph node. After re-penetration of the antigen, there is a rapid proliferation of specifically sensitized lymphocytes, which releases pro-inflammatory substances (lymphokines), causing an inflammatory process at the place where the skin contacts the sensitizing substance. The changes usually concern the surface layers of the skin (epidermis and skin warts) and are characterized by erythema, presence of blisters and exfoliation.

The scope of tests in accordance with:

- »Regulation of the European Parliament and of the Council (WE) No. 1223/2009 dated 30.11.2009 regarding cosmetic products
- » COLIPA Guidelines
- "Product test Guidelines for the Assessment of Human Skin Compatibility 1997"
- "Guidelines for the Evaluation of the Efficacy of Cosmetic Products 2008"

#### Product characteristics

Appearance:

Gelled solution

Color:

Colorless

Fragrance:

Perceptible smell of fragrance composition

Purpose:

Hair cleansing

How to use:

Apply the shampoo to wet hair, massage it in, leave it for 2 minutes, then rinse

thoroughly. In case of contact with eyes, immediately rinse with water.

Packaging

Commercial packaging

#### Selection of Study Participants

# The selection of Study Participants was carried out in accordance with the Research Procedure, taking into account:

»Helsinki Declaration of 1964 (with later additions)

»Current Polish and European legal regulations

»Cosmetics Europe guidelines using the inclusion and exclusion criteria

Twenty volunteers (women and men aged 27 to 59) were selected for the study. Volunteers filled out a detailed survey regarding their lifestyle, current state of health, past illnesses, eating habits, medicines and the use of stimulants. All volunteers selected for the study met the requirements for inclusion in the study and signed consent for conscious participation in the study, and were informed about the purpose of the study, how it was conducted and about possible side effects. 90% of people participating in the study reported a history of adverse reactions caused by the use of certain cosmetics and detergents (including allergic reactions). 10% of people participating in the study did not have hypersensitivity and did not report a tendency to adverse reactions in the interview. The skin at the test site was normal, without lesions.

#### Testing methodology

The study was conducted in accordance with the Research Procedure.

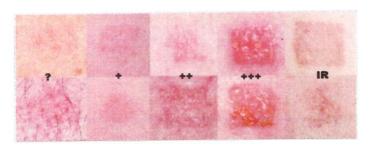
The test substance (10% water solution) is applied to the skin using special chambers fixed on a hypoallergenic adhesive. The chambers are filled with the substances being tested, and then affixed to the skin of the back in the shoulder blade area. During the tests one cannot soak his/her back and should avoid sudden movements and sweating. Patches with test substances are left on the test skin for 48 hours. After removal of the adhesives with the chambers, the excess of test substances is removed by pressing a paper towel onto the skin. The skin reaction is assessed immediately after removing the adhesives, after 72 hours, after 96 hours and after 7 days according to a scale that is consistent with the scale generally accepted in dermatological tests



## **TEST RESULTS**

The patch test readings were made according to the International Contact Dermatitis Research Group (ICDRG)

Graphical scale of the ICDRG patch test reading



( - ) negative reaction

(?) doubtful reaction

(+) weak reaction

( ++ ) strong reaction

(+++) very strong reaction

(IR) irritative reaction

no reaction

subtle erythema, palpably undetectable erythematous spot

palpable erythema, suggestive of mild edema / infiltration, clumping may

occur, no blisters

increased swelling, infiltration, clots, blisters

blisters, erosions, ulceration

shiny skin, dry skin, pimples, erythema

In the group of 20 people who underwent the study, no positive contact allergy or irritant reaction was observed after using the product.

#### SUMMARY

The lack of positive reactions indicates that the tested product does not show irritating and sensitizing effects in contact with the skin.

## Additional information

The issued opinion does not apply to people who are allergic to any of the components of the product under evaluation.

The test sample was prepared by making a 10% water solution specific to rinsed products.

#### THE END of the Attachment

Prepared on: 30-07-2021	Prepared by: JARS Employee no.: 2477	Authorized by: JARS Employee no.: 2427
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## **ANALYTICAL LABORATORIES**

microbiology - physicochemistry - sensory



## TEST REPORT No.: K/0/07/2021/209/F/8/EN

Order No.:

K/0/07/2021/209

- A accredited methodology (AB 1095); reference if the law so provides (the result can be used to assess compliance in the legally regulated area).
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- MON methodology accredited in terms of "OiB"
- GMP+ methodology registered in the scope of GMP+ B11 protocol (feed testing)
  - A/P accredited methodology of the subcontractor
    - P non-accredited methodology of the subcontractor

Materia	al/product tested: Cosmetics							
Produc	t name: PROPAR	IN - Shan	poo Hair	Growth System		Date: 13.	07.2021	
Produce	r:	no	data					
Date of	production:	no	data					
Lot num	iber:	35	2					
Notes of	n the sample:	lac	ck					
-	collected according to: transported by: Shipping				Sample receiver:	JARS employee r	ю.: 2098	
Sample	no.: 12438/07/21 Sample evaluation	n: un	reservedly	y Analysis start da	tte: 13-07-2021 Ana	lysis end date:	14-09-202	1
Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	N
	Application study - panel, subjective	-		PB-34/LK ed. 1 dated 17.12.2020	no requirements	in Attachment		
Ł								

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#### Remarks:

NOTE: The original test reports are issued as PDF file, signed with a qualified electronic signature. Therefore, all prints are copies, unless certified to be true to the original PDF file.

Report prepared in a single copy		The end of the Report	Origin	nal of PDF: Customer, copy of PDF to: Laboratory archive
Created on:	Authorized by:	Approve	ed by:	
27-09-2021	JARS employee no.: 2426	Specjalista ds.l	Kosmetyków	Signed with a qualified electronic signature
		JARS employe	ee no.: 2098	

Measurement uncertainty of the test results is presented when, it is relevant to the validity of applications of the test results applied.

The results relate to the tested samples.

The test report includes test results of the following number of samples: 1 pc(s) and without the written approval of the laboratory shall not be reproduced except in full.

Customer may file complains within 14 days from receiving the report.

The laboratory does not store the samples after testing, unless otherwise agreed with the customer.

Place of performance of the tests (location codes): Ł - Łajski, M - Mysłowice, PS - in situ measurement.



## Attachment No. F/12438/07 to the Study Report

Product name:

PROPARIN - Shampoo Hair Growth System

Sample number:

12438/07/21

Test type:

Application study - panel, subjective

Study start date:

12.08.2021

Study end date:

09.09.2021

#### Aim of the study

The aim of the research was to evaluate and verify the product's performance (properties) declared by the Manufacturer.

#### Methodology

Application tests were carried out on a group of 20 study subjects at home. The tested product was used in accordance with the Manufacturer's directions. After 4 weeks of use, the participants were asked to complete (anonymously) the assessment card accordingly to their own feelings.

#### Suggested usage

Apply the shampoo to wet hair, massage it in, leave it for 2 minutes, then rinse thoroughly. In case of contact with eyes, immediately rinse with water.

## Manufacturer's declarations of the product's properties

The declarations and properties of the product are included in the questions of the questionnaire.

## Characteristics and composition of the group taking part in the study

Total number of people: 20

10% Sex distribution: 90% Male Female 30% 18-30 years old Age distribution: 31-40 years old 35% 20% 41-50 years old 10% 51-60 years old 5% > 61 years old

# RESULTS

Tested feature/property	Possible responses	Number of people
	very high	15%
	high	65%
Efficiency	average	20%
,	low	0%
	very low	0%
	very pleasant	20%
	pleasant	50%
Fragrance	neutral	30%
	unpleasant	0%
	very unpleasant	0%
	proper	100%
Fragrance intensity	too intense	0%
,	not intense enough	0%
	very easy	30%
	easy	70%
Application	average	0%
- Ph.	difficult	0%
	very difficult	0%
	very good	30%
Rinsing	good	70%
	average	0%
<b>g</b>	bad	0%
	very bad	0%
Hair conditioning effect	very clear	0%
	clear	70%
	average	25%
	low	0%
	none	5%
Nourishing effect	very clear	0%
	clear	65%
	average	30%
inouniering enece	low	0%
	none	5%
	very clear	10%
	clear	80%
Hair cleansing effect	average	10%
Tian Gloanienig Gliebe	low	0%
	none	0%
	very high	5%
5000 5000 W W W W	high	50%
Ability of the product to reduce	average	30%
hair loss	low	15%
	none	0%
	very clear	0%
	clear	60%
Effect of thicker hair	average	25%
	low	10%
	none	5%
	very clear	5%
D.	clear	65%
		20%
Effect of healthier hair	average	
Effect of healthier hair	average low	5%



Tested feature/property	Possible responses	Number of people
	very clear	15%
	clear	55%
Improvement of hair hydration	average	10%
, ,	low	15%
	none	5%
	very clear	10%
	clear	55%
Improvement of hair smoothness	average	10%
	low	20%
	none	5%
	very clear	10%
	clear	65%
Improvement of hair softness	average	10%
	low	10%
	none	5%
	very clear	15%
	clear	45%
Improvement of hair elasticity	average	30%
, _	low	5%
	none	5%
	very clear	15%
	clear	60%
Overall improvement of hair	average	15%
condition	low	5%
	none	5%

	Number o	f people [%]
Tested feature/property	NO	TAK
Did you experience any side effects after using the product? Irritation, redness, itching etc.?	100	0

	Number of people [%]			
Tested feature/property	YES	UNDECIDED	NO	
Would you buy the product in the future?	80	5	15	

Tested feature/property	Number of people [%]		
	All right	During illness	After illness
Health status during the course of study	100	0	0

	Number of people [%]		
Tested feature/property	NO	YES	
Did you take any medicine during the during the course of the study?	95	5	



## SUMMARY

PROPARIN - Shampoo Hair Growth System has been evaluated by the study participants positively. The product has a high and very high efficiency (80% of responses) and it has a properly intense (100% of responses), pleasant and very pleasant fragrance (70% of responses). The shampoo has a clear and very clear hair cleansing, nourishing and conditioning effect (80%, 65% and 70% of the responses respectively). Moreover, the product has a clear and very clear effect of healthier (70% of responses) as well as thicker hair (60% of responses). The product clearly and very clearly improved the overall condition of the hair (75% of responses). No one noticed any side effects after using it.

#### THE END of the Attachment

1	Prepared on:	Prepared by:	Authorized by:		
	14-09-2021	JARS Employee no.: 2477	JARS Employee no.: 2426		

