



## Clinical Study Summary Report

# **A Clinical Study for Determining the Sun Protection Factor and UVA Protection Factor of “GLUTANEX UV Glow Balm”**

**Requestor: Nexus Pharma Co., Ltd.**

**February 02, 2024**



**Korea Institute of  
Dermatological Sciences**

# Report Summary



Test Title	A Clinical Study for Determining the Sun Protection Factor and UVA Protection Factor of "GLUTANEX UV Glow Balm"
Research Organization	Korea Institute of Dermatological Sciences 6F, H Business Park Building A, 25 Beobwonro 11-gil, Songpa-gu, Seoul, Republic of Korea
Requestor	Nexus Pharma Co., Ltd. 6F, 71, Gonghang-daero 45-gil, Gangseo-gu, Seoul, Republic of Korea
Test Product	GLUTANEX UV Glow Balm
Formulation	Solid
Test Period	December 26, 2023 ~ February 02, 2024
Methods	<p>This test is conducted in accordance with the JCIA, SPF Test Method of Cosmetics Europe (ISO24444:2019/AMD.2022), ISO24442:2022 and the Standard Operating Procedure (SOP) of Korea Institute of Dermatological Sciences.</p> <ol style="list-style-type: none"><li>1) Selection of subjects: Healthy female and male, aged from 18 to 60 years old</li><li>2) Product Application: Evenly apply <math>2.00 \pm 0.05</math> mg/cm<sup>2</sup> amount to the test area.</li><li>3) Application areas: 35 cm<sup>2</sup> (7 cm x 5cm)</li><li>4) Waiting time after product application: 15-30 minutes</li><li>5) Test device: Multi-port Solar Simulator 601-300W</li><li>6) Assessment Methods<ul style="list-style-type: none"><li>- Evaluate the response of minimal persistent pigment darkening Dose (MPPD) within 2 -24 hours after UV irradiation</li><li>- Evaluate the response of minimal erythral dose (MED) within 16 -24 hours after UV irradiation</li></ul></li><li>7) Statistical criterion: Check whether 95% of confidence interval is within <math>\pm 17\%</math> of mean SPF and UVAPF</li></ol>
Results	<p>The Sun Protection Factor (SPF) and UVA Protection Factor (UVAPF) of "GLUTANEX UV Glow Balm" were SPF <b>60.3 <math>\pm</math> 4.4</b> and PFA <b>17.6 <math>\pm</math> 2.1 (PA++++)</b>, respectively.</p> <p>The skin adverse reaction was not observed during the entire test processes.</p>
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Report Date	February 02, 2024