

**TEST FACILITY**

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**CLIENT**

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<b>Test Report No: MB 15-23922.19</b>	<b>Date: October 16, 2015</b>
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**SAMPLE ID:** The client identified the following test material as “Skin Tite”.

**SAMPLING DETAIL:** Test sample was submitted to the laboratory directly by the client. No special sampling conditions or sample preparation were observed by MB Research Laboratories.

**DATE OF RECEIPT:** Sample was received at MB Research Laboratories facilities on September 10, 2015.

**TESTING PERIOD:** September 18, 2015

**AUTHORIZATION:** Signed project number MB 15-23922.19 signed by Tim Boyer

**TEST REQUESTED:** To predict dermal irritation potential of test articles in the context of identification and classification of skin irritation hazard according to the European Union (EU) classification (R38 or no label), United Nations Globally Harmonized System of Classification and Labeling of Chemicals (GHS) classification system (Category 2 and non-irritants), and OECD Guideline for the Testing of Chemicals No. 439 – In Vitro Skin Irritation: Reconstructed Human Epidermis Test Method. This study is designed based on MatTek protocol *in vitro* EpiDerm™ Skin Irritation Test.

**TEST RESULTS:** The test article submitted has passed and is classified as a non-irritant in accordance with the OECD Guideline for the Testing of Chemicals No. 439.

**Prepared For:**

**Tim Boyer**  
**Technical Director**  
**SMOOTH-ON, INC.**