

**TEST FACILITY**

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**MB Research Laboratories**  
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**CLIENT**

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<b>Test Report No: MB 17-25557.19</b>	<b>Date: December 13, 2017</b>
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**SAMPLE ID:** The client identified the following test materials as “**Derma-tac**” and “**Derma-tac Remover**”.

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

**DATE OF RECEIPT:** Samples were received at MB Research Laboratories facilities on October 26, 2017.

**TESTING PERIOD:** October 30, 2017

**AUTHORIZATION:** Signed project number MB 17-25557.19 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 articles are classified as a non-irritant.

**Prepared For:**

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