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SHENZHEN ESUN INDUSTRIAL CO.,LTD. 15A, MICROSOFT KETONG BUILDING, NO. 55 GAOXINNAN 9TH ROAD, HIGH TECH COMMUNITY, YUEHAI STREET, NANSHAN DISTRICT, SHENZHEN.

Toxicological Risk Assessment (TRA) Cover Page

The following samples were submitted and identified on behalf of the client as:

TPU-95A FILAMENT

(1 formula and 4 additional colors)

Net Weight : 1000G

Item No. : BLACK, BLUE, RED ,GREY, WHITE
Manufacturer : SHENZHEN ESUN INDUSTRIAL CO.,LTD.

Region of Origin : CHINA
Region of Destination : EUROPE

SGS Ref No. : WHIN2412001069PL03

Sample Information Received : FORMULATION WITH CHEMICAL NAME, CAS NO. AND

Order Receiving Date : DEC 26, 2024
Last Information Date : JAN 03, 2025

Assessment Period : DEC 26, 2024 TO JAN 14, 2025

Service Requested

Toxicological Risk Assessment (TRA) for the said product per information provided.

Conclusion

Please refer to the section "CONCLUSION"

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Shenzhen Branch Testing Center Hardlines.

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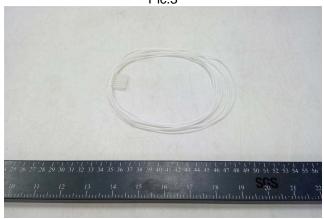
Sample Photo:

Pic.1 Pic.2





Pic.3



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******* For Further Details, Please Refer to the Following Page(s) *******

Signed for and on behalf of SGS-CSTC Standards Technical Services Co., Ltd. Shenzhen Branch Testing Center

Giai

Xue Na, Ella Senior Engineer Scan to see the report

SGS

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Verification:

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TOXICOLOGICAL RISK ASSESSMENT REPORT

The formula of TPU-95A FILAMENT-Black, blue, red, grey, white is reviewed. The product is for EU market and intended for printing as client claims. The net weight of solid (The formula under assessment) is 1000g.

This risk assessment takes account of:

Corrosivity
Skin irritation
Eye irritation
Strong sensitization
Oral toxicity
Inhalation toxicity (if applicable)

According to:

Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures and its following amendments, referred as the CLP Regulation

The followings were considered in this toxicological assessment

Ingredient toxicological profile Potential ingredient interaction Consumer exposure scenario

Section I. FORMULATION REVIEW

According to the client's needs and to protect business secrecy, detailed formulations are NOT presented in this report, whereas they are recorded in the SGS Archives. In case of medical events, it is the client's responsibility to provide the formulations to a patient's physician, or SGS upon request may furnish relevant medical units, on a confidential basis, information necessary to diagnose or treat cases of exposure without prior written authorization.

The formulation is not classified for human health according to the CLP Regulation. None of the ingredients in the formulation is classified as carcinogenic, mutagenic and toxic to reproduction (CMR) Cat. 1A, 1B or 2 under the CLP Regulation.

No allergenic fragrance has been indicated to be present in the formulation.

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Section II. CONSUMER HEALTH RISK ASSESSMENT

Afore-mentioned formulation has been reviewed for safety at the basis of the following assumptions.

Exposure assessment

The following assumptions have been made for assessment of exposure:

- Product category: TPU-95A filament
- Physical form: solid
- Intended use (suggested use): for printing
- Accessibility: this product will contact with the consumer directly
- Exposure route(s): primarily via dermal contact
- Default body weight: 60 kg
- Target population: adult as client claims

Section III. RISK EVALUATION

This product is unlikely to cause damage to internal organs through skin contact in majority of consumers under normal and reasonably foreseeable conditions of use.

Section IV. CONCLUSION

The formulation is not classified as a Dangerous Mixture as per the human health criteria of the CLP Regulation. Provided the manufacturer's instructions are followed, it is considered that, in the present state of knowledge, the submitted formulation put on the market is unlikely to pose a significant risk to the health of intended consumer under normal and reasonably foreseeable conditions of use.

Recommended Safety Labelling

Not Applicable.

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The TRA assessment is valid under the following conditions:

- 1. Potential physical, mechanical, electrical and microbiological risks of the product have not been assessed and evaluated.
- 2. The formulation, description and other supporting documents of the product were submitted and identified by/on behalf of the client, assumed to be valid and accurate.
- It was assumed that either the ingredients, or the finished product contain no contaminants or residues that cause toxicity to a consumer who may be exposed. All polymers should be well-cured if they are present in the formulation.
- 4. The risk evaluation can solely be applied to and appropriate for, the product described above under the stated conditions. Modification of any components of the finished product, such as net weight, adjustment of ingredient concentrations, should therefore require re-evaluation.
- 5. Heavy metal contents in the product were not assessed and evaluated.
- 6. This product has been assessed for compliance only with the regulation(s) specified herein.
- 7. If there is an adverse reaction from using this formulation then the undersigned should be informed so that the formulation can be further reviewed.
- 8. This product has been evaluated for human health only, environmental concern/ ecotoxicity is not assessed in this report.
- The assessment is performed by in-house method, taking reference of available consumer product exposure database, literature data to the best of knowledge by the time of assessment.



HONGJING XU MB, MPH, Diplomate, American Board of Toxicology, European Registered Toxicologist, Certified Toxicologist of CSOT Certified Cosmetic Safety Assessor

*****	End of	Report	*****
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