BIO-TECHNOLOGIES

## DECLARATION OF CONFORMITY

The Company

| Name: | Advanced Bio-Technologies Inc. |
| :--- | :--- |
| Address: | 4830 WEST KENNEDY BLVD, SUITE 600, TAMPA, FL 33609 |

With the EU Authorised Representative:
Name: Alliance Pharma S.r.l
Address: Viale Restelli 5, 20124, Milan, Italy

## DECLARES

Device Name:
Other Trade Names:

Device Classification:

## KELO-COTE UV

Kelo-cote Solaire, Dervida cicatrices gel SPF30
I (Rule 1)

Conformity assessment procedure: Directive 93/42 EEC Annex VII
, meets the essential requirements stated in Annex I of the European Directive $93 / 42 / E E C$ and its amendments Directive 2007/47 EC.
v is put on the market with the CE mark, according to the European Directive 93/42/EEC and its amendments till Directive 2007/47 EC.

- Applicable harmonised standards applied to this device:

| Standard / Guideline <br> Number | Name / Title |
| :--- | :--- |
| BS EN ISO 13485:2012 | Medical Devices- Quality Management Systems - Requirements for Regulatory <br> Purposes |
| BS EN ISO 14971:2012 | Medical Devices- Application of Risk Management to Medical Devices |
| BS EN ISO 10993-1: 2009 | Biological evaluation of medical devices -- Part 1: Evaluation and testing within a <br> risk management process |
| BS EN ISO 10993-5:2009 | Biological Evaluation of Medical Devices-Part 5: Tests for In-vitro cytotoxicity |
| BS EN ISO 10993-10:2009 | Biological Evaluation of Medical Devices-Part 10: Tests for irritation and skin <br> sensitisation |
| BS EN ISO 15223-1:2012 | Medical Devices- Symbols to be used with Medical Device labels, labelling and <br> information to be supplied General Requirements |
| BS EN 1041: 2008 + A1:2013 | Information supplied by Manufacturers with Medical Devices |
| MEDDEV 2.7.1 | Guidelines on Medical Devices; Clinical Evaluation |
| MEDDEV 2.12/1 | Guidelines on Medical Devices; Guidelines on a Medical Devices Vigilance System |
| MEDDEV 2.12/2 | Guidelines on Medical Devices; Post Market Clinical Follow Up Studies |

The manufacturer undertakes to keep, and place at the Authorities disposal, the device technical file for a period of five years from the last date of manufacture of the product.

Date: $12 \operatorname{Sep} 2016$


Advanced Bio-Technologies Inc.
Legal Representative

