EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60138834 0001

Report No.: 12022699 007

Manufacturer:

Aso Philippines, Inc.
4th Street, Block D-5,
Mactan Economic Zone 1,
Lapu-Lapu City, Cebu 6015
Philippines

Products:

Aspects of manufacture concerned with securing and maintaining sterile conditions of adhesive bandage strips and wound dressings

Replaces Approval, Registration No.: DD 60094742 0001

Expiry Date: 2024-04-23

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2019-04-24

Date: 2019-04-24

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle TÜVRheinland

Notified Body

M.Sc. M. Aihara

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.