## Distel

## High-Level Medical Surface Disinfectant CLP Compliance Factsheet

The appearance of Distel Medical products is changing to comply with the new Classification, Labelling and Packaging (CLP) Regulation.

CLP is the European Interpretation of GHS (Global Harmonisation System) for Classification, which is a UN initiative introduced in 1992. Its aim is to drive greater consistency in chemical labelling across the world, with the goal of increasing user and environmental safety associated with use of chemicals.

The CLP Regulation replaces the Dangerous Substances Directive (DSD) and the Chemicals Hazard Information and Packaging for Supply (CHIP) Regulation.

CLP becomes mandatory from 1 June 2015. This means that the appearance of labels and Material Safety Data Sheets of Distel Medical products will change. Already manufactured and labelled products will continue to be supplied after 1 June 2015 until stocks last, resulting in a mix of CHIP and CLP labels during a short period.

Please contact your local Tristel representative or Customer Services to request CLP Material Safety Data Sheets.

For further detailed information regarding the CLP Regulation, please visit www.echa.europa.eu.

## Distel is manufactured in Great Britain by:

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Use biocides safely. Always read the label and product information before use.



