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## **EC DECLARATION OF CONFORMITY**

CellPath Ltd declares that the IVD Medical Device range described hereafter

## 39. Archiving Group of Products

have been classified as Annex III and are in conformity with the Essential Requirements (Annex I) and provisions of Council Directive 98/79/EC,

and are in conformity with the national standards transposing harmonised standards EN ISO 13485:2012, EN 13612:2002, EN 15223-1:2012, ISO 14971:2012,

and are subject to the procedure set out in Annex III of Directive 98/79/EC.

**PLJ Webber** 

Director

On behalf of CellPath Ltd

Date: 24/03/2014

