

EC DECLARATION OF CONFORMITY

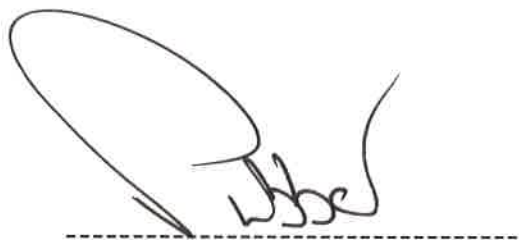
CellPath Ltd declares that the IVD Medical Device range described hereafter

08. Cassettes and Biopsy Handling 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