

Viby J 2022-08-26

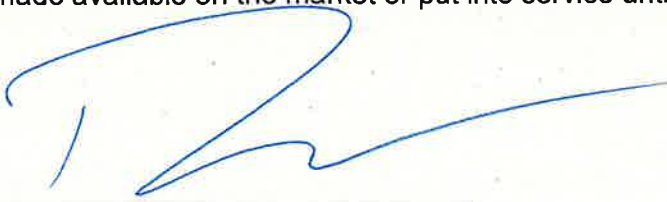
**To whom it may concern**

**Micro well group culture dish, 9-well and 16-well (REF 16604 and 16606 respectively) are CE marked and in conformity with Medical Device Directive (MDD) 93/42/EEC.**

Vitrolife A/S has discontinued the production of Micro well group culture dish, 9-well (REF 16604) and Micro well group culture dish, 16-well (REF 16606).

Last production lots of REF 16604 and REF 16606 were CE marked and released for sale in May 2022 under the EC certificate No. 10000448014-PA-NA-DNK Rev 0.0 (expired June 29 2022 (attached to this letter).

According to article 120 paragraph 4 of the Regulation (EU) 2017/745 – Medical Device Regulation (MDR), medical devices produced by Vitrolife A/S under MDD may continue to be made available on the market or put into service until 26 May 2025.



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QCRA Manager