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In reply please
refer to: UK/DM/QSS/sj

Your reference:

Mr. Jeff Hawkins
Sensitech Inc.
A Business Unit of Carrier Corporation
Suite 258X
800 Cummings Center
Beverly, MA 01915
Etats-Unis d'Amérique

05 June 2009

Dear Mr. Hawkins,

We refer to your application for pre-qualification of your product "VaxAlert™" Type 1 and Type 2, manufactured by Sensitech, USA.

We have now received the results of the device evaluation and are pleased to inform you that your product "VaxAlert™" has been recommended for inclusion in the PQS database of immunization devices pre-qualified for procurement by United Nations Agencies. The product will appear in the WHO web site with the following code:

:: PQS E06/10 VaxAlert TM, Sensitech, USA

We would like to draw again your attention to the terms and conditions and definitions set out in the pre-qualification information pack. Note particularly the requirement that you inform us of any future changes to the device specification or changes to the manufacturing site, and that you notify us of any complaints, product recalls, component defects and other events that may affect the device. Also note that all pre-qualified devices are subjected to annual evaluation which you will be contacted for each annual review by the WHO PQS secretariat.

Yours sincerely,

Mr Denis Maire
PQS Secretariat
Quality, Safety and Standards

cc: Katinka Rosenbom, UNICEF SD
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