**Public Summary**

**Summary for ARTG Entry:** 240813 Integrated Sciences Pty Ltd - OraQuick ADVANCE® Rapid HIV-1/2 Antibody Test and Kit Controls - HIV1/HIV2 antibody IVD, kit, immunochromatographic test (ICT), rapid

**ARTG entry for**
Medical Device Included - IVD Class 4

**Sponsor**
Integrated Sciences Pty Ltd

**Postal Address**
PO Box 731, WILLOUGHBY, NSW, 2068 Australia

**ARTG Start Date**
5/06/2015

**Product category**
Medical Device Class 4

**Status**
Active

**Approval area**
IVD

**Conditions**
- The inclusion of the kind of device in the ARTG is subject to compliance with all conditions placed or imposed on the ARTG entry. Refer Part 4-5, Division 2 (Conditions) of the Therapeutic Goods Act 1989 and Part 5, Division 5.2 (Conditions) of the Therapeutic Goods (Medical Devices) Regulations 2002 for relevant information.
- Breaching conditions of the inclusion related to the device of the kind may lead to suspension or cancellation of the ARTG entry; may be a criminal offence; and civil penalties may apply.

**Manufacturers**

**Name**
Orasure Technologies Inc

**Address**
220 East First Street
Bethlehem, PA, 18015-1360
United States Of America

**Products**

1. **OraQuick ADVANCE® Rapid HIV-1/2 Antibody Test and Kit Controls - HIV1/HIV2 antibody IVD, kit, immunochromatographic test (ICT), rapid**

**Product Type**
IVD

**Effective date**
11/12/2018 11:43:07 AM

**GMDN**
48454 HIV1/HIV2 antibody IVD, kit, immunochromatographic test (ICT), rapid

**Functional description**
The OraQuick ADVANCE® HIV-1/2 Test is a manually performed, visually read, 20 minute immunoassay for the qualitative detection of antibodies to HIV-1 and HIV-2. The assay test strip, contains synthetic peptides representing the HIV envelope region and a goat anti-human IgG procedural control immobilized onto a nitrocellulose membrane in the Test (T) zone and the Control (C) zone, respectively

**Intended purpose**
Intended to detect HIV antibodies in blood, oral fluid and plasma by ICT

**Variant information**

**Specific Conditions**
1. The person (the sponsor) in relation to whom the 'OraQuick ADVANCE Rapid HIV-1/2 Antibody Test and Kit Controls' (the Device) is included in the Australian Register of Therapeutic Goods (the ARTG) must ensure that the Device is only supplied for use by:
   a. laboratories that are accredited by the National Association of Testing Authorities (NATA) as medical testing laboratories and that participate in an HIV point of care quality assurance program; or
   b. organisations that:
      i. will perform, or supervise the performance of, HIV testing using the device; and
      ii. have received training in the delivery and administration of HIV testing in accordance with the requirements of the National HIV Testing Policy; and
      iii. participate in an HIV point of care quality assurance program.

2. The sponsor of the 'OraQuick ADVANCE Rapid HIV-1/2 Antibody Test and Kit Controls' must make available training in the correct use of the Device and interpretation of results.

3. The sponsor must maintain records that demonstrate that the device has been supplied in compliance with condition 1 and that it has complied with condition 2.

4. The sponsor must provide to the Therapeutic Goods Administration (TGA) a post market surveillance report for each reporting period commencing on the date of inclusion of the device in the ARTG and ending at the end of the next 30 June and each twelve (12) months thereafter for the next five (5) financial years. Reports must be provided to the TGA before 1 October after each reporting period and must include the following: a. Numbers of tests sold in Australia and Worldwide
   b. Any adverse events including numbers of any reported false positive or false negative results in Australia and Worldwide.
   c. Reported problems or complaints associated with the use/interpretation of the device in Australia and Worldwide.
   5. Post-market reports must be sent to the TGA at the following email address postmarketdevices@health.gov.au

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This is not an ARTG Certificate document.
The onus is on the reader to verify the current accuracy of the information on the document subsequent to the date shown.