

Ellison Test Certificate



The Ellison Test is the technical audit carried out as part of one of ADISA’s ITAD certifications and is only available to those certified to the ICT Asset Recovery Standard 8 or the ITAD Essentials Certification. The test is carried out at a facility to verify the toolsets used for data sanitisation and includes a forensic assessment of data recovery on a range of storage media. If the test confirms that the product works correctly, is being used properly, and no data was recovered, then the media type is approved (see page 2 of this certificate for details).

Certification Number:	AAC268.ESS	Issue Date:	07.07.2026
Version Number:	v1.0	Expiry Date:	07.07.2027

Testing organisation and forensic analyst name:
ADISA CERTIFICATION LIMITED
JOSHUA SHEPPARD

Pass

British Medical Auctions
UNIT 1, STERLING CENTRE, EASTERN ROAD,
BRACKNELL,
RG12 2PW

The results presented in this document are only valid within the forensic environment that was set up during the test. The full results are in the client report and available at the client's discretion.



To verify the authenticity of this document, scan the QR code.

A handwritten signature in black ink, appearing to read "Sheppard".

Forensic Analyst signature

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Approved Sanitisation Capability

	Reuse	Destruction
<u>Specialist Devices (eg Medical Equipment)</u>	☒	☒

For re-use the applicant shall have a written method for the type of specialist device to achieve the following:

- Storage Identification** - Identify whether each the device contains volatile storage or not.
- Reset Functionality** - Identify what capability the native device user interface provides and whether a reset or other sanitisation function is available.
- Post-reset Accessibility** - If a reset or other sanitisation function is available – perform a test to verify reset effectiveness to ensure that subsequent users of the device cannot access original data/information stored in the device.
- Repeatability** - Create a written sanitisation methodology for each type of specialist device for future use and build up a knowledge bank.
- Audit Validation** - Following each sanitisation process a unique certificate or other form of evidence shall be created.
- Audit Verification** - Quality control shall be carried out by a member of the team who is not involved in the sanitisation process.

NB: The use of software can be added to the method but is not required.

As the appointed forensic analyst for this technical audit, I confirm that British Medical Auctions have evidenced competency to securely process and sanitise the above media and product types.



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Forensic Analyst signature