

Pathlinks (Lincoln)
 HTA licensing number 12314

Licensed under the Human Tissue Act 2004

Licensed activities

The table below shows the activities this establishment is licensed for and the activities currently undertaken at the establishment.

Area	Making of a post-mortem examination	Removal from the body of a deceased person (otherwise than in the course of an anatomical examination or post-mortem examination) of relevant material of which the body consists or which it contains, for use for a scheduled purpose other than transplantation	Storage of the body of a deceased person or relevant material which has come from a human body for use for a scheduled purpose
Hub site Lincoln County Hospital	Licensed	Licensed	Licensed
Mortuary	<i>Carried out</i>	<i>Carried out</i>	<i>Carried out</i>
Pathology lab	-	-	<i>Carried out</i>
Accident and emergency department	-	<i>Carried out</i>	-
Satellite site Pilgrim Hospital	Not Licensed	Licensed	Not Licensed

Mortuary	-	<i>Carried out</i>	-
Accident and emergency department	-	<i>Carried out</i>	-
Satellite site Grantham and District Hospital	Not Licenced	Licensed	Not Licenced
Mortuary	-	<i>Carried out</i>	-
Accident and emergency department	-	<i>Carried out</i>	-

Summary of inspection findings

The HTA found the Designated Individual (DI) and the Licence Holder (LH) to be suitable in accordance with the requirements of the legislation.

Although the HTA found that Pathlinks (Lincoln) ('the establishment') had met the majority of the HTA's standards, two major and three minor shortfalls were found against standards for Governance and quality systems and Premises, facilities, and equipment. These related to maintenance of body store and PM room areas, fridge and freezer capacity and mortuary lone working procedures.

The HTA has assessed the establishment as suitable to be licensed for the activities specified, subject to corrective and preventative actions being implemented to meet the shortfalls identified during the inspection.

Compliance with HTA standards

Major shortfalls

Standard	Inspection findings	Level of shortfall
PFE1 The premises are secure and well maintained and safeguard the dignity of the deceased and the integrity of human tissue		
a) The premises are clean and well maintained	<p>Although the mortuary premises are subject to regular cleaning, deterioration of some areas means that they cannot be cleaned and decontaminated effectively.</p> <p>Examples include:</p> <ul style="list-style-type: none"> • Cracking of the floor tiles in the main PM suite and failure of the grouting. • The bases of the body store fridges have exposed concrete. • There is damage to the walls in the body store and above the dissection benches in the PM room exposing plaster. • Although a leak in the roof of fridge room B has been fixed, it has caused damage to the floor and the surface is lifting 	Major
PFE2 There are appropriate facilities for the storage of bodies and human tissue		
c) Storage for long-term storage of bodies and bariatric bodies is sufficient to meet needs	<p>There is insufficient freezer storage capacity to meet needs.</p> <ul style="list-style-type: none"> • There are four standard size freezer spaces for the long-term storage of bodies over three hospital sites. • There is no bariatric freezer storage. 	Major

Minor Shortfalls

Standard	Inspection findings	Level of shortfall
GQ1 All aspects of the establishment's work are governed by documented policies and procedures		
a) Documented policies and SOPs cover all mortuary/laboratory procedures relevant to the licensed activity, take account of relevant Health and Safety legislation and guidance and, where applicable, reflect guidance from RCPATH.	Lone working is commonplace, however there are no specific lone working procedures in place.	Minor
PFE2 There are appropriate facilities for the storage of bodies and human tissue		
b) There is sufficient capacity for storage of bodies, organs and tissue samples, which take into account predicted peaks of activity	There is a temporary Portacabin unit with 70 spaces that is being used continually. The Trust have plans to remove the unit which will result in the establishment having insufficient capacity for the storage of bodies.	Minor
d) Fridge and freezer units are in good working condition and well maintained.	The fridge and freezer units are very old. The seals on the doors are detaching and there is a build-up of frost within the freezer.	Minor

The HTA requires the DI to submit a completed corrective and preventative action (CAPA) plan setting out how the shortfalls will be addressed, within 14 days of receipt of the final report (refer to Appendix 2 for recommended timeframes within which to complete actions). The HTA will then inform the establishment of the evidence required to demonstrate that the actions agreed in the plan have been completed.

Advice

The HTA advises the DI to consider the following to further improve practice:

Number	Standard	Advice
1.	C1(c)	<p>The establishment do not carry out perinatal or paediatric PMs, however they take consent using the referring hospitals consent documentation.</p> <ul style="list-style-type: none">• The consent form was due for review in February 2022• The consent form does not make it clear who to contact should families wish to withdraw consent. <p>The DI is advised to flag these issues to the partnering Trust.</p>
2.	GQ1(a)	<p>The DI for this licence is not referenced within the '<i>Incident Reporting Guidance</i>' (INT-INS-73) document. The DI is advised to include this information.</p>
3.	GQ1(g)	<p>The establishment is part of a joint Pathology service with other neighboring hospitals and there is some overarching governance relating to licensable activity. The DI may wish to consider meeting with the DI from the other HTA licence to strengthen this harmonised approach.</p>
4.	GQ2(a)	<p>Some of the areas that are covered in the establishments audits refer to the HTA's old standards and standards for the Anatomical Examination sector. Following substantial stakeholder engagement, revised standards came into effect in 2017, and the DI is advised to make sure that the relevant standards are referenced in all future audit activities.</p>
5.	PFE3(a)	<p>The dissection boards within the PM room are starting to show signs of wear and the DI is advised to replace them.</p>

Background

Pathlinks (Lincoln) has been licensed by the HTA since August 2007. This was the fourth inspection of the establishment; the most recent inspection took place in July 2016.

Since the previous inspection, there has been some significant changes to the licence arrangements including the change of Designated

Individual (DI) in October 2017 and a change in Corporate Licence Holder contact in February 2022.

Description of inspection activities undertaken

The HTA's regulatory requirements are set out in Appendix 1. The inspection team covered the following areas during the inspection:

Standards assessed against during inspection

All 72 HTA licensing standards were covered during the inspection (standards published 3 April 2017).

Review of governance documentation

The inspection team reviewed the establishment's self-assessment document provided by the DI and Mortuary Manager in advance of the inspection. Policies and procedural documents relating to licensed activities were reviewed. This included standard operating procedures, risk assessments, audits, incidents, ventilation, and equipment servicing reports, and training and competency assessment documents. Consent seeking procedures and information for relatives giving consent for adult and perinatal PMs was also reviewed.

Visual inspection

The inspection team undertook a visual inspection of the hub and satellite premises which included the mortuary body storage areas, the PM suite as well as the storage arrangements for relevant material held within the facilities.

Audit of records

The inspection team undertook audits of traceability for seven bodies in storage. This included community and hospital cases. Traceability details were crosschecked between the identification band on the body and information in the mortuary register and electronic records. No discrepancies were identified.

Audits were conducted of stored tissue taken at PM examination for five cases. Information was crosschecked between the mortuary documentation, Coroner's paperwork, family wishes forms, the mortuary electronic database, and tissue being stored. No discrepancies were identified.

Meetings with establishment staff

The assessment team met with staff carrying out activities under the licence, including the Mortuary Manager, an Anatomical Pathology Technologists (APT), a senior porter, staff involved in the consent seeking processes, the Bereavement Midwife, and the DI.

Report sent to DI for factual accuracy: 16 May 2022

Report returned from DI: 31 May 2022

Final report issued: 06 June 2022

Completion of corrective and preventative actions (CAPA) plan

Based on information provided, the HTA is satisfied that the establishment has completed the agreed actions in the CAPA plan and in doing so has taken sufficient action to correct all shortfalls addressed in the Inspection Report.

Date: 17 May 2023

Appendix 1: The HTA's regulatory requirements

Prior to the grant of a licence, the HTA must assure itself that the DI is a suitable person to supervise the activity authorised by the licence and that the premises are suitable for the activity.

The statutory duties of the DI are set down in Section 18 of the Human Tissue Act 2004. They are to secure that:

- the other persons to whom the licence applies are suitable persons to participate in the carrying-on of the licensed activity;
- suitable practices are used in the course of carrying on that activity; and
- the conditions of the licence are complied with.

Its programme of inspections to assess compliance with HTA licensing standards is one of the assurance mechanisms used by the HTA.

The HTA developed its licensing standards with input from its stakeholders. They are designed to ensure the safe and ethical use of human tissue and the dignified and respectful treatment of the deceased. They are grouped under four headings:

- consent
- governance and quality systems
- traceability
- premises facilities and equipment.

This is an exception-based report: only those standards that have been assessed as not met are included. Where the HTA determines that there has been a shortfall against a standard, the level of the shortfall is classified as 'Critical', 'Major' or 'Minor' (see Appendix 2: Classification of the level of shortfall). Where HTA standards are fully met, but the HTA has identified an area of practice that could be further improved, advice is provided.

HTA inspection reports are published on the HTA's website.

Appendix 2: Classification of the level of shortfall

Where the HTA determines that a licensing standard is not met, the improvements required will be stated and the level of the shortfall will be classified as 'Critical', 'Major' or 'Minor'. Where the HTA is not presented with evidence that an establishment meets the requirements of an expected standard, it works on the premise that a lack of evidence indicates a shortfall.

The action an establishment will be required to make following the identification of a shortfall is based on the HTA's assessment of risk of harm and/or a breach of the Human Tissue Act 2004 (HT Act) or associated Directions.

1. Critical shortfall:

A shortfall which poses a significant risk to human safety and/or dignity or is a breach of the HT Act or associated Directions

or

A combination of several major shortfalls, none of which is critical on its own, but which together could constitute a critical shortfall and should be explained and reported as such.

A critical shortfall may result in one or more of the following:

- A notice of proposal being issued to revoke the licence
- Some or all of the licensable activity at the establishment ceasing with immediate effect until a corrective action plan is developed, agreed by the HTA and implemented.
- A notice of suspension of licensable activities
- Additional conditions being proposed
- Directions being issued requiring specific action to be taken straightaway

2. Major shortfall:

A non-critical shortfall that:

- poses a risk to human safety and/or dignity, or
- indicates a failure to carry out satisfactory procedures, or
- indicates a breach of the relevant Codes of Practice, the HT Act and other relevant professional and statutory guidelines, or

- has the potential to become a critical shortfall unless addressed

or

A combination of several minor shortfalls, none of which is major on its own, but which, together, could constitute a major shortfall and should be explained and reported as such.

In response to a major shortfall, an establishment is expected to implement corrective and preventative actions within 1-2 months of the issue of the final inspection report. Major shortfalls pose a higher level of risk and therefore a shorter deadline is given, compared to minor shortfalls, to ensure the level of risk is reduced in an appropriate timeframe.

3. Minor shortfall:

A shortfall which cannot be classified as either critical or major, but which indicates a departure from expected standards.

This category of shortfall requires the development of a corrective action plan, the results of which will usually be assessed by the HTA either by desk based review or at the time of the next inspection.

In response to a minor shortfall, an establishment is expected to implement corrective and preventative actions within 3-4 months of the issue of the final inspection report.

Follow up actions

A template corrective and preventative action plan will be sent as a separate Word document with both the draft and final inspection report. Establishments must complete this template and return it to the HTA within 14 days of the issue of the final report.

Based on the level of the shortfall, the HTA will consider the most suitable type of follow-up of the completion of the corrective and preventative action plan. This may include a combination of

- a follow-up inspection
- a request for information that shows completion of actions
- monitoring of the action plan completion
- follow up at next routine inspection.

After an assessment of the proposed action plan establishments will be notified of the follow-up approach the HTA will take.