Histopathologic assessment of cultured human thymus

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Abstract

The maintenance and propagation of complex mixtures of cells in vitro in the form of native organs or engineered organoids has contributed to understanding mechanisms of cell and organ development and function which can be translated into therapeutic benefits. For example, allogeneic cultured postnatal human thymus tissue has been shown to support production of naïve recipient T cells when transplanted into patients with complete DiGeorge anomaly and other genetic defects that result in congenital lack of a thymus. Patients receiving such transplants typically exhibit reversal of their immunodeficiency and normalization of their peripheral blood T cell receptor V-beta repertoire, with long-term survival. This study was designed to assess the histopathologic changes that occur in postnatal human thymus slices when cultured according to protocols used for transplanted tissues. Results showed that as thymic organ cultures progressed from days 0 through 21, slices developed increasing amounts of necrosis, increasing condensation of thymic epithelium, and decreasing numbers of residual T cells. The architecture of the thymic epithelial network remained generally well-preserved throughout the 21 days of culture, with focal expression of cytokeratin 14, a putative biomarker of thymic epithelial cells with long-term organ-repopulating potential. All organ slices derived from the same donor thymus closely resembled one another, with minor differences in size, shape, and relative content of cortex versus medulla. Similarly, slices derived from different donors showed similar histopathologic characteristics when examined at the same culture time point. Taken together, these results demonstrate that diagnostic criteria based on structural features of the tissue identifiable via hematoxylin and eosin staining and cytokeratin immunohistochemistry can be used to evaluate the quality of slices transplanted into patients with congenital athymia.

Introduction

Patients with complete DiGeorge anomaly and other forms of congenital athymia have severe immunodeficiency due to their lack of a thymus and the resulting absence of T-cell production. Transplantation of cultured postnatal thymus tissue has been shown to successfully
and Precision Pathology Center (BRPC; https://pathology.duke.edu/core-facilities-services/biorepository-precision-pathology-center; BRPC@duke.edu). Requestors may be asked to bear costs for scanning slides and/or anonymization of existing scans based on the specific cases requested.

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**Competing interests:** Cultured postnatal human thymus (RVT-802) is an investigational product implanted into patients under an Investigational New Drug (IND) application with the United States Food and Drug Administration (FDA) that is sponsored by Dr. Markert. The technology for RVT-802 was developed by Dr. Markert and has been licensed to Enzyvant Therapeutics GmbH (Enzyvant) by Duke University. Dr. Markert and Duke University have received royalties from Enzyvant. If the technology is commercially successful in the future, Dr. Markert and Duke University may benefit financially. Portions of the salaries of Drs. Markert, Kurtzberg, and Hale and consultation fees for Dr. Neff were paid by funding to Duke University from Enzyvant. This does not alter our adherence to PLOS ONE policies on sharing data and materials.

Support production of naïve T cells in patients with congenital athymia due to DiGeorge anomaly and other genetic causes (e.g. deficiency of TBX2 or FOXN1) [1–8]. Transplanted patients develop naïve peripheral blood T cells with normalization of the peripheral blood T cell receptor V-beta repertoire and immune function [5,9]. Biopsy of thymus tissue 2–4 months after transplantation demonstrates colonization of donor thymus stroma by recipient thymocytes, with normal-appearing cortex and medulla and Hassall bodies [10,11]. Overall survival in transplanted patients is 72% (43/60) at 1 year and 73% (36/49) at 2 years. In contrast, almost all untreated patients die by age 2 due to overwhelming infection [5,7,12,13]. The mechanism by which thymic transplantation restores immune function in these patients is hypothesized to be due to signaling stimulated by direct contact of thymic epithelial cells in the transplanted thymus with recipient T cell precursors that leads to generation of mature functional T cells.

Thymus tissue intended for transplantation is obtained with permission from the parent(s) of immunocompetent infant donors undergoing cardiac surgery. The donated tissue then undergoes a series of processing steps that include slicing and culture for 12 to 21 days [5,11]. The purpose of culture is to partially deplete T cells from the thymic epithelial network. This depletion provides space for colonization of the depleted thymus slices by recipient T cell precursors and also minimizes the potential for graft-versus-host disease mediated by donor T cells. This study describes the histopathologic changes that occur during the culture of thymus slices. Understanding these changes can potentially lead to the validation of enhanced histopathologic criteria for prospective assessment of the quality of cultured thymus slices prior to transplantation, based on characteristics of tissues that have successfully generated immune reconstitution in prior recipients [12].

**Materials and methods**

Thymic tissue was obtained from immunocompetent infant donors <9 months of age who were undergoing corrective cardiac surgery where removal of a portion of the thymus was routinely required to facilitate the cardiac repair. The parent(s) of each donor provided written informed consent to allow any thymic tissue that was removed and otherwise would be discarded to be potentially used for transplantation or research. These studies were approved by the Institutional Review Board of Duke University Medical Center. The donor thymus used for the research portion of this study was sliced and cultured in a Good Manufacturing Process (GMP)-compliant cell manufacturing laboratory using donor qualification and culture procedures identical to those used for thymus samples intended for transplantation [2,3,5,14,15]. However, rather than transplantation into recipients, all slices from these research lots were fixed in 10% neutral buffered formalin at specific time points during culture and entirely submitted for histopathologic examination. All slices from 5 thymuses were submitted at a single time point at either day 0 (56 slices), day 5 (23 slices), day 9 (28 slices), day 12 (21 slices), or 21 (12 slices). For the remaining 3 thymuses, 1/5 of the slices representing that thymus were submitted for examination at each of days 0, 5–7, 9, 12, or 20–21 (n = 28 total slices per time point). Fixed slices were submitted to a Clinical Laboratory Improvement Act (CLIA)-certified, College of American Pathologists (CAP)-accredited clinical laboratory, where formalin-fixed paraffin embedded blocks, hematoxylin and eosin (H&E)-stained sections, and an immunohistochemical panel were prepared using procedures identical to those used to assess lots intended for transplantation. Antibodies used included pan-cytokeratin (clones AE1/AE3; Leica), cytokeratin (CK) 14 (clone LL02; Leica), CD3 (clone LN10; Leica), and Ki-67 (clone MIB-1; Dako). Automated immunohistochemistry was performed using standard immunoperoxidase methodologies and 3, 3′-diaminobenzidine (brown) substrate, with a hematoxylin...
counterstain. Some of these slices were also evaluated for reactivity with antibodies specific for terminal deoxynucleotidyl transferase (clone TdT; clones SEN28; Leica), CD45RO (clone UCHL1; Dako), CD43 (clone DF-T1; Dako), and the T cell receptor β chain (βF1, clone 8A3, Thermo Scientific). H&E and immunohistochemical slides (AE1/AE3, CK14, CD3, Ki-67) were also examined from 11 lots of cultured thymus that were implanted into athymic recipients contemporaneously and up to 2.5 years before preparation of the research lots described above.

Results

Initial identification of tissue as thymus

The thymus is a bi-lobed organ that is present in the anterior mediastinum. It possesses a thin connective tissue capsule that is typically removed during processing within the laboratory. However, extensions of the capsule (trabeculae) and associated blood vessels, fibrous tissue, adipose tissue, and variable numbers of mature hematopoietic cells may still be observed in thymus slices. The trabeculae divide the thymus tissue into lobules that are composed of a lacy-appearing three-dimensional network of thymic epithelial cells with intervening spaces containing the developing T cells (thymocytes). The lobules typically demonstrate an outer cortex and an inner medulla, which vary in their histologic appearance as well as phenotype and function of the cells they contain. The thymic cortex is very basophilic (blue) due to densely packed immature thymocytes that stain very darkly with hematoxylin. The density of the more mature thymocytes that are present in the thymic medulla is lower than that of the cortex, so the medulla tends to appear more eosinophilic (pink). The thymic medulla also contains pathognomonic eosinophilic cytokeratin-containing structures called Hassall bodies. Macrophages present within the thymus may form “tingible bodies” that appear as a light-colored circular area against a background of darkly staining thymocytes (a “starry sky” pattern). Large numbers of tingible bodies are characteristic of stress involution, which may occur in normal thymus donors due to the stress of severe cardiac defects, surgery, and/or corticosteroid treatment and does not disqualify a thymus for transplantation. The typical histologic features of sliced normal thymus prior to culture (Day 0) are shown in Fig 1.

Assessment of Hassall bodies

The thymic medulla contains characteristic structures called Hassall bodies that are composed of terminally differentiated thymic epithelial cells that react with monoclonal antibodies that also react with the terminally differentiated upper layers of the epidermis (reviewed in [16]). Hassall bodies may serve as a marker of tissue identity, since they are found only in thymus. They may also reflect the quality of the input donor tissue, since the terminal differentiation process is normally triggered by thymocyte-thymic epithelial cell interactions [16]. Hassall bodies are usually readily identified by their appearance as eosinophilic whorls of epithelial cells on H&E-stained slides (Fig 2). The central-most layers typically lack nuclei. Variable amounts of cellular debris may also be present in the center of the Hassall bodies. Hassall bodies stain very strongly with AE1/AE3 antibodies, which can be used as secondary confirmation of the identity of these structures if not definitively identified in H&E-stained slides.

General thymocyte changes during culture

Thymocytes are progressively lost as thymus tissue is cultured, either from being flushed out during media changes or from thymocyte death followed by their degradation within the tissue (Figs 3–6). However, different from what occurs in vivo, dead cells may persist in cultured
thymus long-term due to inability to recruit phagocytes to clear them. Thymocyte nuclei undergoing cell death may show pyknosis (chromatin condensation) and karyorrhexis (nuclear fragmentation) but typically, as these cells deplete their energy but are not phagocytosed, they show ragged nuclear edges with loss of nuclear membrane integrity. Karyolysis (complete dissolution of nuclei in necrotic cells) typically occurs within 2–3 days in vivo, but appears to occur more slowly during thymus culture. After 5–9 days in culture, most thymocytes have retained nuclei, but nuclear membranes are not intact, indicating early necrosis (Fig 4). These altered nuclear characteristics facilitate the distinction between viable and non-viable thymocytes, even when nuclei are retained. Large foci of eosinophilic necrotic cell debris where thymocyte nuclei have undergone karyolysis may also be seen (Fig 4A).

**Histologic assessment of cell integrity and tissue architecture**

When cultured thymus tissue is to be transplanted, it is critical to be able to accurately distinguish between slices with extensive thymocyte necrosis but thriving thymic epithelial cells (the optimal tissue for transplantation) versus slices with similar amounts of necrosis where thymic epithelial cells are also compromised. Although histologic examination of standard H&E slides cannot directly assess viability, it can assess the degree of preservation of normal structures and the presence or absence of indicators of cell death. As described above, thymocytes are expected to be depleted during the culture process. Many thymocytes are washed from the slice during the daily media changes. Many others will die in situ, where their nuclei and cell bodies may remain for prolonged periods of time. However, the membrane integrity of dead thymocytes is compromised and their nuclei exhibit “ragged” edges. Thymocyte debris may clump together, making it impossible to discern individual cell borders. Thymocyte death and

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**Fig 1. Architecture of cultured thymus, day 0.** A. B. Hematoxyl in and eosin (H&E) staining shows well-defined cortical and lighter-staining medullary areas, as expected for normal pediatric thymus. C. Immunohistochemistry with a cocktail of antibodies (AE1/AE3) that together detect all types of epithelial cells demonstrates that thymic epithelial cells are present beneath the capsule and in a light lacy network in both cortex and medulla. Arrows in B and C point to a Hassall body. D. Cytokeratin 14 (CK14) antibody reacts with thymic epithelial cells in the sub-capsular cortex and in the medulla, as well as with scattered thymic epithelial cells in the cortex. The dotted line highlights an area of medulla that is surrounded by cortex. Brown color indicates a positive reaction with antibody. SCC denotes sub-capsular cortex, Cor denotes cortex, and M denotes medulla. Bar represents 1 mm in A and 500 μm in B–D.

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depletion is an expected and desirable consequence of culture and relative lack of intact thymocyte nuclei is potentially reflective of slice quality. The thymic epithelial cells are more easily visualized as thymocytes are depleted from the tissue. The nuclei of viable thymic epithelial cells are typically oval, larger than those of thymocytes, and have a sharply defined nuclear membrane outlined by the hematoxylin stain, with one or more nucleoli. These thymic epithelial nuclei typically look “open”, meaning they do not stain darkly with hematoxylin. This fits with an interpretation that they are alive and metabolically active, since active chromatin (“euchromatin”) does not bind the hematoxylin dye. The presence of nucleoli, which are the sites of ribosome synthesis, in many thymic epithelial cells further confirms that they were alive and metabolically active at the time of fixation. Examples of intact and viable-appearing thymic epithelial cells are shown in Fig 7.

Careful microscopic examination of multiple thymus slices at any given time point from day 0–21 showed that all slices derived from the same donor thymus closely resembled one another. The differences observed between different slices derived from the same thymus as a function of culture time were primarily related to the amount of necrosis (increased as culture time increased) and numbers of residual thymocytes (decreased as culture time increased). Slices derived from different donors were also qualitatively similar to each other when examined at similar time points. Differences between different donor tissues included relative size, shape, relative content of thymus versus medulla (but both were always present on each slice examined), amount of necrosis, condensation of thymic epithelium, and numbers of residual

Fig 2. Examples of Hassall bodies. The histologic appearance of Hassall bodies is shown on day 0 (A, B) and day 9 (C, D) of culture. Hematoxylin and eosin (H&E) stain is shown in panels A and C. Reactivity with pan-cytokeratin (AE1/AE3) antibodies is shown in panels B and D; brown color indicates a positive reaction. Arrowheads point out representative Hassall bodies, which appear less prominent on H&E-stained sections of cultured thymus due to the depletion and necrosis of the surrounding thymocytes. However, Hassall bodies can still be readily identified by careful examination or by using immunohistochemistry. Bar represents 100 μm.

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Fig 3. Architecture of cultured thymus, day 7. A. B. Hematoxylin and eosin (H&E) stain shows marked depletion of thymocytes, although some cortical areas (Cor) still contain large numbers of thymocytes with retained nuclei. Pan-cytokeratin (AE1/AE3) (C) and cytokeratin 14 (CK14) immunohistochemistry (D) show condensation of the thymic epithelium in the subcapsular cortex (SCC) and in the medulla (M). Brown color indicates a positive reaction with antibody. Bar represents 1 mm in A and 500 μm in B–D.

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Fig 4. Architecture of cultured thymus, day 9. A. B. Hematoxylin and eosin (H&E) stain. Few if any live T cells or thymic epithelial cells are present in the pale-staining area in A that is enclosed by the dotted line, which is almost completely necrotic (Necr). Most nuclei formerly present in this region have been degraded, a process called karyolysis. Other areas where the nuclei from residual thymocytes have not been completely degraded continue to stain dark blue with hematoxylin. Arrow in B points to a Hassall body. C. Pan-cytokeratin (AE1/AE3) immunoreactivity. D. Cytokeratin 14 (CK14) immunoreactivity. Brown color indicates a positive reaction with antibody. Bar represents 1 mm in A and 500 μm in B–D.

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T cells. Most changes in histologic appearance had already occurred by day 5 of culture, with generally only additional depletion of thymocytes as cultures progressed.

Assessment of thymic epithelial architecture

A cocktail containing anti-cytokeratin antibodies AE1 and AE3 detects essentially all thymic epithelial cells. In contrast, an antibody reactive with CK14 detects a subset of thymic epithelial cells that are present in the subcapsular cortex and seemingly scattered throughout the remainder of the cortex and medulla. Some of these CK14+ cells have been suggested to have the potential to differentiate into both cortical and medullary epithelial cells [17] and thus to represent long-term repopulating cells. As thymocytes are depleted from the tissue, the thymic epithelial cells become more visible. The three-dimensional thymic epithelial network is normally demonstrated in sections via a light and lacy arrangement of connected epithelial cells and/or seemingly scattered thymic epithelial cells whose connections are not evident in the section being examined. The three-dimensional thymic epithelial network contracts to varying degrees as thymocytes are lost during culture. This can result in condensation of the residual epithelium, such that the subcapsular cortical epithelial layer becomes thicker and medullary thymic epithelial cells become more tightly packed. Examples of these changes are shown in Figs 1 and 3–6 for a single thymus lot examined on days 0, 7, 9, 12, and 20. Despite the loss of numerous thymocytes and potentially large areas containing necrotic debris, the overall architecture of the thymic epithelial cell network remains generally well-preserved through at least 21 days of culture (Fig 8).

Assessment of residual T cells

As described above, the primary purpose of culture of donor thymus tissues that are intended for transplantation is to partially deplete T cells to facilitate colonization of the transplanted

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**Fig 5. Architecture of cultured thymus, day 12.** A, B. Hematoxylin and eosin (H&E) stain. At this time point, many thymocytes have either been lost from the tissue or they have died and their nuclei have been dissolved, making the tissue more eosinophilic (pink). Some areas retain architecture characteristic of normal uncultured thymus with cortical-like areas (Cor) that stain more basophilic (blue) and medullary-like areas (M), although with greatly decreased thymocyte cellularity. C. Pan-cytokeratin (AE1/AE3) immunoreactivity. D. Cytokeratin 14 (CK14) immunoreactivity. Brown color indicates a positive reaction with antibody. At this time point, the sub-capsular cortex (SCC) has thickened and epithelial cells appear more prominent due to the decreased numbers of thymocytes present. Arrows point to representative Hassall bodies. Bar represents 1 mm in A and 500 μm in B–D.

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slices with recipient thymocyte precursors and to decrease the risk of graft-versus-host disease. Immature T cells (thymocytes) and mature T cells can be identified in histologic sections by their morphology as well as via immunohistochemical stains that detect proteins specifically expressed by these cell types. CD3 is a component of the T cell receptor for antigen that is present on >95% of cortical and medullary thymocytes as well as all mature T cells. On day 0, the plasma membranes of essentially all immature T cells in the cortex and the medulla appear strongly reactive with CD3 antibody in a membrane pattern (Fig 9A and 9B). Viable thymocytes/T cells continue to react with CD3 antibody in a membrane pattern as cultures continue. However, the CD3 antibody also reacts strongly with dead thymocytes, including the anucleate debris that remains after dead thymocytes undergo karyolysis (Fig 9D, 9G, 9I and 9K). So much thymocyte debris remains that slices may still appear strongly and uniformly brown when viewed at low magnification (Fig 9C, 9F, 9H and 9J), although examination of the slide at higher power (e.g. 40X objective) confirms that relatively few intact potentially viable thymocytes are present at these time points. Non-viable cells and cellular debris will be cleared by recipient phagocytes after transplantation and therefore convey no risk of graft-versus-host disease. It is not generally possible to accurately identify CD3+ membrane immunoreactivity in tissue areas where considerable amounts of strongly staining debris prevents assessment of cellular membranes of adjacent cells. However, even at late time points during culture, CD3 immunohistochemistry typically highlights at least some thymocytes/T cells that have intact plasma membranes suggesting that they may still be viable (Fig 9E, 9G, 9I and 9K).

To determine whether other thymocyte-reactive antibodies that were currently validated by our laboratory for clinical diagnostic use could better identify residual immature or mature T cells in cultured thymus tissues, patterns of immunohistochemical reactivity were qualitatively

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**Fig 6. Architecture of cultured thymus, day 20.** A, B. Hematoxylin and eosin (H&E) stain. At this time point, most thymocytes have either been lost from the tissue or they have died and their nuclei have been dissolved, making the tissue more eosinophilic (pink). Large groups of residual thymocytes are rare, although scattered cells with nuclear characteristics of thymocytes are evident. C. Pan-cytokeratin (AE1/AE3) immunoreactivity. Much of the epithelium present in formerly medullary areas (M) is condensed due to loss of medullary thymocytes, but scattered epithelial cells indicative of a residual light, lacy, three-dimensional network of thymic epithelial cells remain. D. Cytokeratin 14 (CK14) immunohistochemistry highlights former medullary areas and the subcapsular cortex. Brown color indicates a positive reaction with antibody. Arrows point to representative Hassall bodies. Bar represents 1 mm in A and 500 μm in B–D.

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assessed for cultured thymus reacted with antibodies that recognized CD45RO, terminal deoxynucleotidyl transferase (TdT), CD43, and the beta chain of the T cell receptor (βF1). As shown in Figs 10–13, CD45RO antibody had a similar pattern of reactivity with cortical versus medullary thymocytes as CD3 antibody, while fewer overall thymocytes were reactive with TdT, CD43, and βF1 antibodies. However, TdT, CD45RO, CD43, and βF1 antibodies also reacted with thymocyte debris, and use of these antibodies did not improve identification of the small numbers of intact-appearing residual thymocytes late in culture.

Fig 7. Assessing intact nuclei in thymus slices. Examples of intact thymic epithelial cell nuclei (arrows) are shown in the subcapsular cortex on day 9 (panel A) and in the medulla on day 21 (panel B). Hematoxylin and eosin stain; bar represents 50 μm.

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Assessment of cellular proliferation

The Ki-67 antigen is expressed in the nucleus of all cells that are proliferating (i.e. not in the G0 phase of the cell cycle). On day 0, the majority of the immature thymocytes present within the thymic cortex are proliferating and their nuclei react strongly with antibody specific for the Ki-67 proliferation antigen. In contrast, only rare more mature medullary thymocytes and/or epithelial or stromal cells react with this antibody (Fig 14A and 14B). The pattern of immuno-reactivity observed after day 1–2 of culture is of scattered rare positive cells, almost all of which have the larger nuclei characteristic of thymic epithelial cells (Fig 14C–14K). Thus, Ki-67 reactivity may be a useful adjunct to nuclear appearance by H&E for documenting the viability of thymic epithelial cells within the slice.

Validation of histopathologic assessment criteria using clinical thymus lots

From 1995 through 2019, qualification of cultured thymus lots for transplantation has simply required documentation of sterility (lack of bacterial and mycoplasma contamination) and fulfillment of simple qualitative histopathologic diagnostic criteria that identify the tissue as thymus and document its viability and tissue organization (Tables 1 and 2). Careful microscopic examination of the 8 research lots submitted in toto for histologic examination showed that different slices derived from the same or different thymus tissue were qualitatively similar and were ranked identically per Tables 1 or 2 at any given time point in culture. Differences between slices derived from the same donor thymus were limited to relative size, shape,
content of cortex versus medulla (but both were always present on each slice examined),
amount of necrosis, condensation of thymic epithelium, and numbers of residual T cells. Fur-
thermore, since the major change that occurred between days 5–9 and 12–21 was progressive
depletion of thymocytes, the tissue histology on day 5–9 was reflective of that observed on days
12–21, just with more thymocyte depletion at the later time points.

Retrospective review was performed for 11 cultured thymus lots that were processed either
contemporaneously with or up to 2.5 years prior to these research lots then transplanted into
athymic recipients. All 11 lots met the diagnostic criteria shown in Tables 1 and 2
for slices examined on culture days 0, 5–10, and 12–21. The histologic appearance of each of these slices
(n = 33) closely resembled that of the research lots examined at a similar time point, with no
substantive deviations from the detailed descriptions above. Of note, all 11 recipients devel-
oped sufficient T cell reconstitution to resist infections, with naive CD4 T numbers ranging
from 74 per mm$^3$ to normal for age, when tested 1–3 years post-implantation of cultured
thymus.

Discussion

This study describes histopathologic changes that occur when postnatal human thymus is cul-
tured for up to 21 days. As thymic cultures progressed, slices developed increasing amounts
of necrosis, increasing condensation of thymic epithelium, and decreasing numbers of residual T
cells. The thymic epithelial network remained intact throughout the 21 days of culture, with
continued expression of cytokeratin 14, a putative biomarker of thymic epithelial cells with
long-term organ-repopulating potential. Slices from the same thymus were qualitatively simi-
lar, such that a single slice could adequately represent the entire thymus. Variability in histo-
logic appearance of cultured thymus slices derived from different donors was also minimal at
given any time point. The tissue histology observed early during culture (e.g. days 5–9) closely
reflected what was observed later in culture (e.g. days 12–21), although more thymocyte deple-
tion and necrosis were observed at the later time points.

Immunohistochemistry for antibodies that recognize CK14 or all types of cytokeratins
(AE1/AE3) was helpful in evaluation of cultured thymus slices. Cytokeratin intermediate fila-
ments are important components of the cytoskeleton of all epithelial cells. The pan-cytokeratin
AE1/AE3 stain demonstrates the epithelial network within the examined thymus slice, which
may not be easily discernable at later time points using H&E stain alone. The specific type of
cytokeratin expressed by a particular thymic epithelial cell has been shown to depend on its
stage of development and differentiation and functional state. In mice, antibodies that react
with CK5, CK8, and CK14 have been most commonly used to identify thymic epithelial cell
subtypes [17]. Expression of CK14 was determined in this study, based on previous studies
where CK14 expression was hypothesized to be a characteristic of thymic epithelial cells with
the potential to differentiate toward either cortical or medullary lineages [18]. While the
Fig 10. CD45RO immunoreactivity in thymus slices as a function of time in culture. A, B. On day 0, essentially all immature T cells in the cortex (Cor) and medulla (M) react strongly with antibody specific for CD45RO. Higher magnification (B) shows the strong brown positive membrane reactivity. C, D. By day 7, reactivity with CD45RO antibody is strongest in the medulla. Some positive cells exhibit the expected specific membrane staining, but there is also considerable reactivity with cellular debris. Loss of membrane immunostaining and intense reactivity primarily with residual thymocyte debris is seen on days 9 (E, F), 12 (G, H) and 21 (I, J), although some cells with intact membrane staining can occasionally still be detected (arrows in H). Bar represents 500 μm in panels A, C, E, G, and I and 50 μm in B, D, F, H, and J.

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Clinical outcome of long-lasting immune reconstitution [5] suggests that thymic epithelial progenitors are likely present within transplanted thymus slices, the precise phenotype for such progenitors has not been definitively identified in humans. The phenotype of thymic epithelial progenitors in mice is still controversial [19–22] and only low numbers have been detected.
using multiple antibodies and multi-color flow cytometry. Thus direct detection of thymic epithelial progenitors in human tissue sections is not currently feasible.

These studies identified potentially large amounts of necrotic cellular material remaining within thymic slices at culture days 12–21, the time points when the tissue might be
transplanted. Biopsy examination several months after transplantation shows no evidence of this necrotic material [11]. We hypothesize that the initial presence of this necrotic debris and its underlying extracellular matrix at the time of transplant may serve to preserve the

Fig 12. CD43 immunoreactivity in thymus slices as a function of time in culture. A, B. On day 0, immature thymocytes in the cortex (Cor) react weakly and those in the medulla (M) react more strongly with antibody specific for CD43. The dark bands in A are sectioning artifacts (wrinkles and folds in the tissue). Higher magnification (B) shows the brown positive membrane reactivity with both cortical and medullary thymocytes. C, D. By day 7, reactivity with CD43 antibody is generally limited to the medullary thymocytes. Some positive cells exhibit the expected specific membrane staining, but there is also considerable reactivity with cellular debris in cortical areas. Loss of membrane immunostaining and intense reactivity primarily with residual thymocyte debris is seen on days 9 (E, F), 12 (G, H) and 21 (I, J), although some cells with intact membrane staining can be detected at these later time points. Bar represents 500 µm in panels A, C, E, G, and I and 50 µm in B, D, F, H, and J.

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functional architecture of the slices, including the cortical and medullary niches for developing thymocytes. Experimental use of thymus decellularization approaches that preserve extracellular matrix to recreate murine thymic “organoids” that can support cellular differentiation of both epithelial cells and hematopoietic precursors [23] supports this hypothesis.

Fig 13. βF1 immunostaining in thymus slices as a function of time in culture. A, B. On day 0, a subset of immature T cells in the cortex (Cor) and essentially all of the medullary thymocytes react strongly with βF1 antibody that is specific for the β chain of the αβ T cell receptor. Higher magnification (B) shows strong brown positive membrane reactivity; M denotes medulla. C, D. Reactivity with antibody remains strong on day 7, however the specific membrane staining is lost and reactivity is primarily with cellular debris in medullary areas. Similar loss of membrane immunostaining and reactivity primarily with residual thymocyte debris is seen on days 9 (E, F), 12 (G, H) and 21 (I, J), but cells with intact membranes can be detected (see arrows in J). Bar represents 500 um in panels A, C, E, G, and I and 50 um in B, D, F, H, and J.

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Previous studies demonstrated that the growth potential of thymic epithelial cells was robustly maintained under the culture conditions used, such that cytokeratin-positive epithelial monolayers could be established from the slices up to 12 weeks after initiation of organ culture [10]. The current study did not directly assess epithelial cell viability, but instead used the presence of intact nuclei as a surrogate marker. Since the histopathologic examination focused on evaluation of cellular and tissue structures that could be readily discerned by staining with H&E or reactivity with cytokeratin antibodies, it was inherently limited by the inability of these stains to directly measure viability and cellular function. Thus, histopathologic examination using only H&E stains combined with immunohistochemistry for cytokeratins cannot necessarily address all factors that may influence the quality of the cultured thymus slices. A positive reaction of thymic epithelial cell nuclei with antibodies that recognize the Ki-67 proliferation antigen provides proof that those cells are alive, since Ki-67 is only expressed in actively proliferating cells, i.e. not in G0 phase. However, the sensitivity of this approach is likely limited by the relatively small percentage of thymic epithelial cells that are proliferating at any given point in time.

We also note that, based on our studies to date, it is not possible to link overall patient survival or the time required for their immunoreconstitution with specific histopathologic characteristics of the specific lot of cultured thymus received. Viable-appearing thymus allograft has been identified post-mortem in at least one patient who died post-transplantation [11]. Patient factors such as severe congenital malformations or infections can lead to death prior to T cell reconstitution. Furthermore, clinical requirements for allograft-damaging therapy such as prolonged high-dose corticosteroids may prevent function, independent of thymus lot quality. Studies to identify additional biomarkers that track with thymic epithelial viability and potential function are on-going and will help to definitively address the issue of thymus allograft quality at implantation versus adverse clinical condition in governing the survival and immune reconstitution of transplanted patients.

In summary, histopathologic examination using H&E and cytokeratin immunoreactivity can be useful for assessment of cultured thymus slices. The value of the histopathologic examination to prospectively assess the quality of slices intended for transplantation into patients with congenital athymia may potentially be improved by identifying and developing assays for new biomarkers that reflect thymic epithelial cell viability and/or function.

Table 1. Diagnostic criteria for thymus submitted on day 0

| Assay      | Acceptance Criteria                                                                 | Result Required for Acceptance |
|------------|-------------------------------------------------------------------------------------|---------------------------------|
| Identity   | 50% of sampled tissue contains keratin AE1/AE3 networks in lacy staining pattern     | Present                         |
| Potency    | Hassall bodies identified; CK14 staining in lacy pattern                             | Present                         |
| Viability  | >90% intact nuclei observed                                                        | Present                         |

*Any additional immunostains performed (e.g. CD3 or Ki-67) should also be commented upon.

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Table 2. Diagnostic criteria for thymus that has been cultured for one or more days*.

| Assay         | Acceptance Criteria                                      | Result Required for Acceptance |
|---------------|----------------------------------------------------------|--------------------------------|
| Identity      | Areas positive for keratin AE1/AE3                        | Present                        |
| Potency       | At least one Hassall body identified; CK14 staining scattered throughout | Present                        |
| Viability     | Intact thymic epithelial nuclei observed                  | Present                        |

*Any additional immunostains performed (e.g. CD3 or Ki-67) should also be commented upon.

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