Dynamics of thymus organogenesis and colonization in early human development

Alison M. Farley1,*, Lucy X. Morris1,‡, Eric Vroegindeweij3,‡, Marianne L. G. Depreter1,‡, Harsh Vaidya1, Frances H. Stenhouse1, Simon R. Tomlinson1, Richard A. Anderson2, Tom Cupedo3, Jan J. Cornelissen3 and C. Clare Blackburn1,§

SUMMARY
The thymus is the central site of T-cell development and thus is of fundamental importance to the immune system, but little information exists regarding molecular regulation of thymus development in humans. Here we demonstrate, via spatial and temporal expression analyses, that the genetic mechanisms known to regulate mouse thymus organogenesis are conserved in humans. In addition, we provide molecular evidence that the human thymic epithelium derives solely from the third pharyngeal pouch, as in the mouse, in contrast to previous suggestions. Finally, we define the timing of onset of hematopoietic cell colonization and epithelial cell differentiation in the human thymic primordium, showing, unexpectedly, that the first colonizing hematopoietic cells are CD45+CD34int/–. Collectively, our data provide essential information for translation of principles established in the mouse to the human, and are of particular relevance to development of improved strategies for enhancing immune reconstitution in patients.

KEY WORDS: Thymus development, Human, Hematopoietic, Epithelium, Mesenchyme, Colonization

INTRODUCTION
T-cell development occurs in the thymus and depends on a diverse array of functionally distinct epithelial cell types within the thymic stroma. These thymic epithelial cell (TEC) types originate from the endoderm of the third (and in some species fourth) pharyngeal pouches (PPs), transient bilateral endodermal structures that generate both the thymus and parathyroid glands (Le Douarin and Joteureau, 1975; Gordon et al., 2004). The molecular and cellular processes that govern thymus organogenesis are best understood in the mouse (Blackburn and Manley, 2004; Manley and Condie, 2010). By contrast, knowledge of human thymic organogenesis is based primarily on histological studies, with molecular insight currently limited to analysis of genetic abnormalities affecting thymus development; principally, DiGeorge Syndrome, the human nulde syndrome and APECED, which are caused by mutations in TBX1, FOXN1 and AIRE, respectively (Frank et al., 1999; Baldini, 2005; Villaseñor et al., 2005).

The human thymus is thought to develop from either the third PP (Norris, 1938) or from both the third and fourth PP (Van Dyke, 1941). As in the mouse, the bilateral human primordia initially each comprise thymic and parathyroid domains, and are encased in a neural crest-derived mesenchymal capsule from early in development. From week 7 [Carnegie stage (CS) 18-19] to mid-week 8 (CS20-21) of gestation the thymic component of this primordium migrates ventrally, such that the leading tip of the migrating thymic rudiment remains attached to the parathyroid by a thin, elongated, highly lobulated, cord-like structure. This structure then resolves, separating the parathyroid and thymic primordia (Norris, 1938), and the thymic primordium meet and attach at the pericardium by mid-week 8 (Norris, 1938). It is well documented that an accessory thymus can exist in the cervical region in humans (Norris, 1938; Van Dyke, 1941) and, similarly, the occurrence of cervical thymi in the mouse has recently been reported (Dooley et al., 2006; Terszowski et al., 2006).

The early human thymic primordium is proposed to contain undifferentiated epithelial cells, as reported for the mouse (Lampert and Ritter, 1988; Bennett et al., 2002; Rossi et al., 2006). Medullary development has been observed from week 8 and distinct cortical and medullary compartments by week 16. The presence of other intrathymic cell types, including mesenchymal, vascular and lymphoid cells, has been reported from mid-week 8 (Haynes et al., 1984; Haynes and Heinly, 1995). Between 14 and 16 weeks, mature lymphocytes begin to leave the thymus to seed the peripheral immune system (Van Dyke, 1941; Lobach and Haynes, 1987). However, although the morphological events in human thymus organogenesis are well documented and some information exists regarding TEC differentiation, molecular regulation of these processes remains poorly understood. In particular, only very limited information exists regarding the extent to which mechanisms known to govern thymus development and function in the mouse are conserved in human. Furthermore, neither the time of onset of TEC differentiation nor the identity of markers defining thymic epithelial progenitor cells (TEPCs) has been determined in the human fetal thymus. Similarly, neither the timing of initial hematopoietic colonization nor the identity of the colonizing cells has been reported. Improved understanding is urgently required, owing to the clinical need for improved strategies for enhancing or replacing thymus function in patients (Wils and Cornelissen, 2005; Lynch et al., 2009; Awong et al., 2010).

1MRC Centre for Regenerative Medicine, Institute for Stem Cell Research, School of Biological Sciences, University of Edinburgh, CRM Building, 5 Little France Drive, Edinburgh EH16 4UL. 2MRC Centre for Reproductive Health, Queen’s Medical Research Centre, University of Edinburgh, Edinburgh EH16 4JL. 3Erasmus University Medical Center, Department of Hematology, 3000 CA Rotterdam, The Netherlands.

*Present address: Melbourne Brain Centre, Department of Anatomy and Neuroscience, The University of Melbourne, Parkville, 3010, Australia
‡These authors contributed equally to this work
§Author for correspondence (c.blackburn@ed.ac.uk)

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial Share Alike License (http://creativecommons.org/licenses/by-nc-sa/3.0), which permits unrestricted non-commercial use, distribution and reproduction in any medium provided that the original work is properly cited and all further distributions of the work or adaptation are subject to the same Creative Commons License terms.
Here, we demonstrate that the major mechanisms currently known to underpin thymus organogenesis are conserved between mouse and human. We further define the age at which overt TEC differentiation is initiated in human fetal thymus, and establish the timing of expression of known mediators of TEC function, including the autoimmune regulator AIRE and the chemokines CCL21 and CCL25. These observations delineate the stage in thymus development at which the thymus is composed principally of undifferentiated thymic epithelial progenitor cells. Finally, we determine the stage of thymus organogenesis at which colonization by hematopoietic, endothelial and mesenchymal cells first occurs, identifying the first colonizing hematopoietic cells as CD45^+CD7^+CD34^-^ and the first thymus-seeding T-lineage progenitors as CD45^+CD7^-CD34^int^, in contrast to expectations from previous studies (Haddad et al., 2006).

These data, to our knowledge, constitute the first detailed analysis of human thymus organogenesis and TEC development that is not based solely on morphological and histological data, and thus provide essential information for the translation of principles established in the mouse to the human.

**MATERIALS AND METHODS**

**Human tissue**

First and second trimester human fetuses were obtained following elective medical termination of pregnancy, and all were morphologically normal. Ethical approval for use of human fetal tissue in these studies was granted by the Lothian Research Ethics Committee or the Ethics Committee of the Erasmus University Medical Center. Consent was obtained from all women in writing, and the tissue was anonymized before being made available for research. All experiments using human tissue were performed at the University of Edinburgh or the Erasmus University Medical Center. Embryos were aged according to the standard head/rump measurement and Carnegie stage was determined (Gasser, 1975; O’Rahilly and Muller, 1987). Embryos used in this study were from mid- to late week 6 (CS17), week 7 (CS18-19), early to mid-week 8 (CS20-21), and weeks 9, 10, 15-16 and 17, with days/weeks indicating days/weeks post-fertilization. Note that Carnegie stage does not apply to second trimester fetuses.

**In situ hybridization**

Whole-mount *in situ* hybridizations were performed as previously described (Gordon et al., 2001). Probes were made by PCR amplification from human fetal tissue using primers as listed in supplementary material Table S1.

**Immunohistochemistry**

Immunohistochemistry was performed as previously described (Gordon et al., 2004). Primary antibodies used were: anti-cytokeratin (rabbit polyclonal anti-keratin, Dako); anti-CLDN4 (3E2C1, Zymed); anti-human CD45 (clone H130, BD Pharmingen); anti-keratin 5 (A1F138 rabbit IgG, Covance); anti-keratin 8 (CAM 5.2, Becton Dickinson); anti-keratin 14 (AF64, Covance); TE7 (Developmental Studies Hybridoma Bank); UE1 (biontinated, Vector Labs); anti-AIRE (D-17, Santa Cruz); anti-CD205 (MG38, Serotec); CDR2 (provided by B. Kyewski, German Cancer Research Center, Heidelberg, Germany); anti-HLA-DR/DP/DQ (clone TU39, BD Pharmingen); anti-EPCAM (Ber-EP4, DakoCytomation); anti-CCL21 (anti-6Ckine, R&D Systems); anti-CCL25/TECK (R&D Systems); mouse anti-CD11b/Mac1 (ICRF44, BD Pharmingen); mouse anti-CD3 (WM59, BD Pharmingen); mouse anti-CD144 PE (16B1, eBiosciences); mouse anti-CD34 FITC (clone 581, BD Pharmingen); mouse anti-CD3 Alexa700, mouse anti-CD45 PerCP-Cy5, mouse anti-HLA-DR APC-Cy7, mouse anti-CD123 PE (all BD Pharmingen). Unconjugated antibodies were detected with goat anti-mouse Alexa647 and goat anti-mouse Alexa 568 (Invitrogen).

**RESULTS**

**Spatial and temporal expression of essential regulators of mouse thymus organogenesis is conserved in human thymus development**

To determine whether molecular mechanisms that regulate thymus development are conserved between mouse and human, we investigated the detailed spatial and temporal expression profiles of known regulators of mouse thymus organogenesis in human embryos. We first confirmed the relative staging of human and mouse thymus development. The earliest human tissue available for this study was early week 6 (CS16). At this stage, the third PP had formed and morphological criteria indicated that third PP/thymus primordium development was at an equivalent stage to that at embryonic day (E)10.5 in CBAxC57BL/6 F1 mouse embryos. By early week 7 (CS18), outgrowth of the third PP to form the common thymus/parathyroid primordium (Gordon et al., 2001) had occurred, equivalent to ~E11.5 in the mouse (Fig. 1A). Migration of the thymus component of the primordium along the carotid artery was observed by mid-week 8 (CS21), as described, and at this and subsequent stages the thymus primordium was surrounded by a

**Flow cytometry**

Thymic single cell suspensions were generated by using enzymatic dissociation as described for mouse fetal thymus (Nowell et al., 2011) or by mincing thymic fragments through a 100-μm filter (BD). Flow cytometric analysis was performed on a LSRII Flow Cytometer (BD) and analyzed using FlowJo software (Tree Star). Antibodies used were: mouse anti-CD19 DY590 (Exbio Praha), mouse anti-BDCA2 APC (Miltenyi Biotech), rat anti-CD7 (MCA344, Serotec), mouse anti-human CD45 (clone H130), mouse anti-human CD34 FITC (clone 581), mouse anti-CD3 Alexa700, mouse anti-CD45 PerCP-Cy5, mouse anti-HLA-DR APC-Cy7, mouse anti-CD123 PE (all BD Pharmingen). Unconjugated antibodies were detected with goat anti-mouse Alexa647 and goat anti-mouse Alexa 568 (Invitrogen).

**Grafted re-aggregate fetal thymic organ culture**

Two human thymic lobes (early week 8) were lightly dissociated as described, mixed with 150,000 mouse primary embryonic fibroblasts and re-aggregated as described previously (Sheridan et al., 2009). The re-aggregates were cultured on a filter for 24 hours and then grafted under the kidney capsule of NOD/SCID recipient mice the next morning.rafts were recovered after 2 weeks for analysis.
mesenchymal capsule (Fig. 1B-B'). By week 10, the thymic primordium comprised numerous lobules (Fig. 1C,C'; note that the mid-week 8 thymus is not lobulated), and lobules could also be seen within the elongated cord connecting the thymus and parathyroid primordia (Fig. 1C,C'). Of note is that from week 8, the mesenchyme surrounding the human thymic primordium was less closely associated with the primordium than is the mesenchymal capsule in the mouse (Fig. 1C, arrow).

We then investigated the expression pattern in week 6 human embryos of Tbx1, Hoxa3, Pax1 and Pax9 genes, which encode proteins with genetically defined roles in patterning of the pharyngeal endoderm (PE) and/or formation of the third PP. Tbx1 is required for development of the mouse foregut endoderm, where it regulates segmentation of the PE and PP patterning (Xu et al., 2005). Conserved stage-specificity and localization of Tbx1 expression between mouse and human embryos was demonstrated (ISH; Fig. 2A,A'). In the early week 6 (CS16) human embryo, Tbx1 expression was detected in the core mesenchyme of the first, second and third pharyngeal arches (PAs) and in the third PP endoderm (Fig. 2A,A'). Expression was also detected in the epithelium of the caudal half of the otic vesicle (Fig. 2A).

Hoxa3 is expressed in the mouse from E9.5 in the third pharyngeal cleft ectoderm, in the third and fourth PA neural crest cells, and in the third PP endoderm, and is required for thymus and parathyroid formation (Chisaka and Capecchi, 1991; Manley and Capecchi, 1995). Hoxa3 expression was also conserved between mouse and human; the anterior boundary of Hoxa3 expression in the week 6 (CS16-17) human embryo was the third PA and, as in the E10.5 mouse embryo, expression was observed throughout the third PP and surrounding mesenchyme (Fig. 2B,B').

In the mouse, both Pax1 and Pax9 are expressed in the foregut endoderm from E8.5 (Neubüser et al., 1995; Wallin et al., 1996) and are then expressed in all four PPs and in the thymus throughout organogenesis, becoming restricted to subpopulations of TECs
postnatally (Neubüser et al., 1995; Wallin et al., 1996). Here, PAX1 expression was investigated in a week 6 human embryo; expression was observed in all four PPs (Fig. 2C,C′). PAX9 expression was also examined in human embryos at week 6; expression was evident in the third and fourth PPs but was not detected in the first and second pouches (Fig. 2D,D′). Thus, whereas the expression pattern of PAX1 is conserved between mouse and human, PAX9 expression in the human PE is more restricted than in the mouse at the equivalent developmental stage.

Foxn1 is required cell-autonomously for development of mouse TECs but is not required for initiation of thymus organogenesis (Blackburn et al., 1996; Nehls et al., 1996; Newell et al., 2011). Null mutations in Foxn1 result in athymia and hairlessness in mice, rats and humans (Flanagan, 1966; Festing et al., 1978; Nehls et al., 1994; Frank et al., 1999). In the mouse, Foxn1 expression initiates at the ventral tip of the third PP at E11.25 (in CBAxC57BL/6 embryos) and spreads to the entire thymus domain by E11.5 (Gordon et al., 2001). We could not detect FOXN1 expression in the third PP of early week 6 (CS16) human embryos (not shown) but observed strong expression throughout the thymus domain of the third PP by mid-week 6 (CS16-17; Fig. 2E), suggesting that timing of initiation of high-level FOXN1 expression occurs at equivalent developmental stages in human and mouse. FOXN1 expression was also seen throughout the thymic lobes at week 8 (CS20-21; Fig. 2F′). The human thymus has been suggested to develop from both the third and fourth PPs (Norris, 1938; Van Dyke, 1941). However, we did not detect FOXN1 expression in the fourth PP at any stage of development (Fig. 2E′,E′). This establishes that in the human, as in the mouse, the thymus arises solely from the third PP, indicating that the mechanisms of early thymus organogenesis are closely conserved between these species. Of note is that GCM2, which delineates the parathyroid domain within the pouch, is expressed in both the third and fourth PPs in human (Liu et al., 2010), in keeping with the proposition that human parathyroids derive from both the third and fourth PPs.

We also determined the expression pattern of P63 (TP63 – Human Gene Nomenclature Database), because in the mouse p63 (Trp63 – Mouse Genome Informatics) has a pivotal role in many stratified epithelia, including the thymus (Candi et al., 2007; Senoo et al., 2007). In the early week 6 (CS16-17) human embryo, P63 was expressed by the ectoderm surrounding the pharyngeal arches (Fig. 2F), consistent with its expression in the pharyngeal ectoderm in the mouse from E9.5 (Mills et al., 1999). By mid-week 6, P63 expression was observed in the endoderm of the third PP (Fig. 2F′) and by late week 8, it appeared to be expressed by all TECs (Fig. 2F″). Thus, P63 expression in the pharyngeal region is also conserved between mouse and human.

Finally, we analyzed the expression of BMP2 and BMP4, because BMP signaling has been implicated both in thymus development (Gordon et al., 2010) and as a possible regulator of Foxn1 (Tsai et al., 2003; Senoo et al., 2007). BMP4 expression was observed in the second and third PP endoderm and adjacent mesenchyme in the mid-week 6 (CS16-17) human embryo (Fig. 2G,G′), consistent with its expression pattern at E10.5 in the mouse (Patel et al., 2006), whereas BMP2 was expressed in the third and fourth PP endoderm of the early week 6 (CS16) embryo (Fig. 2H), as observed in the mouse from E9.5 (Lyons et al., 1990).

Collectively, these data demonstrate that the spatial and temporal expression of transcription factors and signaling molecules involved in mouse thymus organogenesis is highly conserved in human thymus development, albeit with some minor regional differences. This strongly suggests that the genetic mechanisms controlling thymus development are conserved between mouse and human. Our previous studies have shown that Rhox4 and Plet1 (1600029D21Rik – Mouse Genome Informatics), specific markers of the mouse third PP/early thymus primordium, are not expressed in this region during human thymus organogenesis (Morris et al., 2006; Depreter et al., 2008). Thus, our data indicate that although the expression patterns of all genes implicated in thymus organogenesis by functional studies are conserved between mouse and human, not all gene expression patterns in this region are conserved between species.

**Onset and progression of epithelial cell differentiation in the human thymic primordium**

We next investigated whether patterning of the human thymic primordium into cortical and medullary compartments occurred in a similar manner to in the mouse, and at what stage the onset of TEC differentiation occurred. In the adult mouse thymus, the structural proteins cytokeratin 8 (K8; Krt8 – Mouse Genome Informatics) and keratin 5 (K5; Krt5 – Mouse Genome Informatics) are broadly restricted to cortical TECs (cTECs) and medullary TECs (mTECs), respectively, and additionally are co-expressed by a minor TEC population found at the cortico-medullary junction and scattered throughout the cortex (Klug et al., 1998). These markers are also co-expressed by most TECs in the E11.5 mouse thymic primordium (Bennett et al., 2002), by E12.5, high level K5 expression marks a medullary thymic epithelial progenitor cell population that also expresses high levels of claudin 4 (Cldn4) and, from ~E13.5, co-stains with the mTEC marker lectin UEA1 (Hamazaki et al., 2007).

At early week 7 (CS18), K5 and K8 were co-expressed by most, if not all, TECs in the human thymic primordium, consistent with the expression pattern seen at the equivalent stage (E11.5) in mouse (Fig. 3A-A″). By mid-week 8 (CS21), these markers had begun to segregate such that, although most TEC expressed K8, K5 was more highly expressed by cells along the midline of the thymic lobe and in some scattered cells (Fig. 3B-B″). Expression of EPCAM, a pan-epithelial marker in the mouse thymus that is expressed at higher levels in mouse mTECs than cTECs (Farr and Anderson, 1985), was also similar in the mouse and human thymus. EPCAM was expressed by all TECs in the week 7 human thymic primordium (Fig. 3C-C″) and by week 15 was expressed strongly by mTECs and weakly by cTECs (Fig. 3D). These data suggested that the onset of overt TEC differentiation in human thymus development occurs around mid-week 8 (CS21), demonstrating that, as in mouse, human TEC development is initiated after the initiation of FOXN1 expression (Newell et al., 2011). Consistent with this conclusion, expression of a functional marker of TEC maturation, major histocompatibility complex (MHC) Class II (HLA-DR/DP/DQ), expression of which is FOXN1-dependent in the early mouse thymic primordium (Nowell et al., 2011), was not detected in human TECs until mid-week 8 (Fig. 3E-E″).

We next determined the pattern of UEA1 staining in human thymus development, because in the mouse UEA1 binding identifies mTECs in the thymic primordium from E13.5 (Hamazaki et al., 2007) and, subsequently, most postnatal mTECs (Farr and Anderson, 1985). Surprisingly, as no UEA1 staining is observed at the equivalent stage in the mouse, UEA1 bound most, if not all, TECs within the week 7 human thymus (Fig. 4A-A″). By mid-week 8, UEA1 staining on TEC was restricted to the central area of the thymic lobes (Fig. 4B-B″), suggesting that medullary epithelial progenitor cells had arisen by mid-week 8, and a subpopulation of mTECs binding UEA1 at high intensity were present by week 15 of human fetal development (Fig. 4D-D″). Of note is that some non-epithelial UEA1+ cells were also observed in the human fetal
thymus, unlike in the mouse. These cells were initially seen lining week 8 thymic lobes (Fig. 4B-B’). We also investigated the expression of CLDN4, as the Cldn4+UEA1+ population in the fetal mouse thymus contains progenitors that give rise to Aire+ mTECs (Hamazaki et al., 2007). By week 8, CLDN4 was expressed by TECs in the central area of the thymic lobes (Fig. 2I; Fig. 4E-G’). These CLDN4+ cells also expressed MHC class II (HLA-DR/DP/DQ; Fig. 4H-H’), suggesting that, as in the mouse (Hamazaki et al., 2007), these are the first cells to differentiate into mature medullary TECs. CLDN4 expression remained restricted to a subpopulation of mTECs and, by week 18, identified both small clusters of nucleated mTECs and swirls of cells surrounding an enucleated region that were strongly suggestive of Hassall’s corpuscles (Fig. 4G-G’).

The autoimmune regulator Aire is essential for induction of central tolerance in mice and humans (Mathis and Benoist, 2009) and therefore we analyzed AIRE expression during human thymus development. AIRE+ TECs were not present in the week 10 human thymic primordium (not shown), but were observed in medullary areas by week 13 (Fig. 4I-I’), and had increased in number by week 17 (Fig. 4J-K’).

To investigate differentiation towards the cTEC fate, we determined the expression pattern of CD205 (LY75 – Human Gene Nomenclature Database) (Shakib et al., 2009) and CDR2 (Rouse et al., 1988). CD205 has recently been shown to mark the onset of cTEC differentiation in the mouse and is expressed as early as E12.5 in mouse thymus development (Shakib et al., 2009); later in fetal development, it may mark a common TEPC (Baik et al., 2013). CDR2 is restricted to cTECs in the human adult thymus (Rouse et al., 1988). We were unable to detect CD205 or CDR2 expression in the early or mid-week 7 thymic primordium by immunohistochemistry (Fig. 4L-L’). However by mid-week 8, extensive expression of both markers was observed throughout the thymic lobes (Fig. 4M-M’). CDR2+ TECs within the mid-week 8 thymus did not express K5, suggesting that they may represent cortical sub-lineage-restricted epithelial progenitor cells. Again, these data are consistent with the onset of TEC differentiation occurring around mid-week 8 in the human thymic primordium.

The chemokine CCL21, which plays a role in hematopoietic colonization of the thymus in mice (Liu et al., 2006), was detected in scattered cells in the human thymic primordium at week 11 (Fig. 4P-P’). The number of CCL21+ TECs was substantially increased by week 17 and appeared to be largely restricted to medullary regions (Fig. 4Q-Q’), consistent with its localization in the mouse (Kwan and Killeen, 2004). CCL25, a chemokine expressed by mouse TECs in a Foxn1-dependent manner that also plays a role in hematopoietic colonization of the thymus (Liu et al., 2006; Nowell et al., 2011), was first detected at mid-week 8 (Fig. 4R-S’). CCL25 expression was initially detected in scattered cells throughout the human thymic primordium. By week 17, CCL25 expression was detected only in cortical TECs (Fig. 4U-W’), as shown by co-staining with CDR2 and EPCAM (Fig. 4V-W’), which mark cortical and medullary regions, respectively, by immunohistochemistry (Farr and Anderson, 1985; Rouse et al., 1988).

**Immigration of mesenchymal cells and endothelial progenitors into the human thymic primordium**

Neural crest-derived mesenchymal cells form the capsule and trabeculae of the fetal thymus and also generate the intrathymic pericytes (Foster et al., 2008). It is well established that thymic mesenchyme is at least partly responsible for regulating expansion (Auerbach, 1960; Jenkinson et al., 2003; Anderson et al., 2006) and migration (Griffith et al., 2009; Foster et al., 2010) of the early thymic primordium. However, the timing of the initial immigration of mesenchymal cells into the human thymic primordium has not been determined. TE7 is a marker of human thymic mesenchymal cells that has been reported to stain the mesenchymal cells surrounding the thymic lobes at weeks 7 and 10, and to mark interlobular fibrous septae, vessels and thymic fibrous capsule at week 15 (Haynes et al., 1984). At early week 7, we observed cells stained with TE7 surrounding the thymic primordium, but no stained cells were seen within the thymic epithelium (Fig. 5A-A’). By mid-week 7, formation of trabeculae was apparent and TE7 expression was also seen in the anterior end of the thymic/parathyroid primordium, where it appeared to be in the luminal spaces (Fig. 5B-B’). However, mesenchymal cells were not evident within the thymic epithelium region itself until mid-week 8.
Fig. 4. Expression of markers of cortical and medullary TECs in human thymus development. (A-K) Medullary TEC differentiation. Images show staining with UEA1 (A-D), anti-CLDN4 and anti-HLA-DR/DP/DQ (MHC II) (E-H) and anti-AIRE (I-K). (L-O) Cortical TEC differentiation. Images show staining with CD205 (L-M) and CDR2 (N-O). (P-W) Chemokine expression. Images show staining with anti-CCL21 (P-Q) and anti-CCL25 (R-W). Developmental stages are as shown. Anti-pan-Cytokeratin (PANK) co-stain reveals the entire thymus primordium. Images are representative of at least two independent analyses. Wk, week. Scale bars: 150 μm in A-B, D-F, H-H, N-W; 75 μm in C-C; 47.6 μm in G-G; 50 μm in I-I, L-M; 55 μm in J-K.
(Fig. 5C,C'). At subsequent stages, TE7+ thymic mesenchyme divided the thymus into discrete lobules, and TE7+ cells were also present throughout the epithelial regions (Fig. 5D-E') (Haynes et al., 1984). Thus, the timing of onset of migration of mesenchymal cells into the human thymic primordium closely follows the onset of FOXN1 expression (Fig. 2E), consistent with the timing of this event in the mouse (Foster et al., 2008).

Immigration of vascular progenitors into the mouse thymus is closely linked to that of neural crest-derived cells, and both of these processes are Foxn1 dependent (Mori et al., 2010; Nowell et al., 2011). However, again, the timing of vascular endothelial progenitor entry into the human thymus has not been determined. Our analysis identified the presence of CD34hiCD31+ cells in the connective tissue between the carotid artery and the thymus primordium at mid-week 7 (Fig. 6A–C'); occasional cells at this stage were found within the epithelial component of the primordium (Fig. 6D–D'). By late week 7 to mid-week 8, CD34hi cells were found within the thymic lobes (Fig. 6E–F') and all co-expressed both CD31 (PECAM1 – Human Gene Nomenclature Database) and VEcadherin (CDH5 – Human Gene Nomenclature Database), identifying them as endothelial progenitors (Fig. 6G–G'). Of note is that the CD34 hi cells present within the thymic primordium at late week 7 to early week 8 did not express CD45 (PTRC – Human Gene Nomenclature Database) (Fig. 6E–E'). Blood vessels per se were present by mid-week 9 (note long blood vessel-like structures at mid-week 9) (Fig. 6H–I').

Some variation in timing of colonization with endothelial progenitors was observed: in the thymus primordium from a CS19 fetus (mid to late week 7; 48-51 days old) and a CS20 fetus (late week 7 to early week 8), several CD34hi cells were present within the epithelial compartment, and were localized to one discrete region (Fig. 6D–E'). However, in a CS21 fetus (mid to late week 8; 53-54 days old) CD34hi cells were present in the mesenchymal capsule but were not detected within the epithelial component of the thymus (Fig. 6F–F'). These differences in kinetics of colonization between different fetal thymus are likely to reflect the effects of genetic variation within the human population on fetal thymus development; in this context, it is well established that thymus organogenesis proceeds at different rates in different inbred mouse strains.

**CD45^+CD7^+ T-lineage progenitors are first present at week 8 and lack high level CD34 expression**

Hematopoietic colonization of the human fetal thymus has previously been proposed to occur during early week 8, and the...
week 7 human primordium was reported as being devoid of hematopoietic cells (Haynes and Heinly, 1995). However, as acknowledged by the authors, this study was compromised by the limited material available (Haynes and Heinly, 1995) and, thus, although the timing at which subsequent stages in T-cell development are first observed has been determined (Res and Spits, 1999), the stage at which the earliest hematopoietic cells colonize the human thymus remains unknown. We therefore sought to determine the kinetics of hematopoietic colonization of the human thymus using markers associated with human hematopoietic progenitor cells, including early T-lineage precursors. Of note is that the earliest T-lineage restricted precursors in human are thought to...
be bone marrow-derived CD45+CD34hiCD7+ cells based on functional analysis of hematopoietic cells at week 8-9 of fetal development, the earliest time-point examined (Haddad et al., 2006).

We observed CD45+ hematopoietic cells in the perithymic space at early week 7 (Fig. 7A,A'), and by early to mid-week 8 we detected occasional CD45+ cells within the thymic primordium of some embryos (Fig. 7B,B', green arrow). By early to mid-week 8, CD45+ cells lined the perithymic space and were also distributed throughout the thymic lobes, and by week 10 the numbers of CD45+ cells within the thymus had increased dramatically (Fig. 7C,D). However, no CD34+CD45+ cells could be detected prior to week 9 (Fig. 7E-M'). Co-staining with additional markers indicated that at mid-week 8 a proportion of the CD45+ cells were Mac1+ (IGTAM+) macrophages (Fig. 7F-F'). Of note is that comparison of immunohistochemistry and flow cytometric staining data indicated that only cells expressing high levels of CD34 were detected by immunohistochemistry.

Surprisingly, given the absence of CD34 hiCD45+ cells at this stage, CD45+CD7+ cells were observed within the thymic lobes from early week 8, when they comprised a minor subpopulation of intrathymic CD45+ cells (Fig. 7G-I'; CD45+CD7+Mac1'; CD45+CD7 Mac1' and CD45+CD7 Mac1- cells were all present at this stage). Consistent with previous reports that CD7 identifies extra-thymic hematopoietic precursor cells with T-lineage potential (Haynes and Heinly, 1995; Haynes et al., 2000), we observed some CD7int cells located in the perithymic space. These cells were sometimes juxtaposed to CD7hi cells that appeared to be actively entering the thymus, suggesting that CD7 upregulation might occur upon migration into the thymic epithelium (Fig. 7H-I'). However, intrathymic CD34hiCD7+ cells were not detected until mid-week 9, when they were observed closely associated with the blood vessels (Fig. 7K-J'). Most intra-thymic CD7+ cells at early week 8 were CD34hi, consistent with the presence of CD45hi cells in the fetal liver (FL) (Fig. 7P,Q). A minor population of CD34lo/−CD7+ cells were also present at week 8, which might represent the CD34+/−CD7−CD56+ natural killer (NK) cell population previously described in the FL (Phillips et al., 1992; Bárcena et al., 1993). However, the week 8 hematopoietic population contained T-cell precursors, as culture of week 8 thymic lobes microdissected free of any other tissues resulted in a robust expansion of CD45+ thymocytes (supplementary material Fig. S1). Furthermore, expression of c-Kit (KIT – Human Gene Nomenclature Database), IL7R, pre Ta (PTCRA – Human Gene Nomenclature Database), RAG1, CD3δ (CD3D – Human Gene Nomenclature Database), and TCR β and γ (TRB and TRG – Human Gene Nomenclature Database) constant region mRNA was detected in microarray analysis of whole mid-week 8 fetal human thymic lobes (supplementary material Table S2). Collectively, these data pinpoint the stage of human thymus development at which colonization by hematopoietic progenitors occurs, and identify a novel subpopulation of CD34hiCD7+ cells that might represent intrathymic T lineage-restricted progenitors.

Subsequent analyses confirmed previous studies indicating the appearance of CD4 and CD8 single positive and γδ TCR+ thymocytes, and CD4+CD25+ T regulatory cells in the human fetal thymus by week 15 (Lobach and Haynes, 1987; Galy et al., 1993; Haynes and Heinly, 1995; Cupedo et al., 2005) (data not shown). Low numbers of conventional dendritic cells (cDCs) were detected at week 15, consistent with a previous report indicating their presence at week 12 (Janossy et al., 1986). CD11c+CD11b+ cDCs increased proportionally by week 16 (not shown), together with a numerical increase in the entire cDC population. Of note is that plasmacytoid dendritic cells were already evident at week 15 (Fig. 7V).

**DISCUSSION**

The data presented above address the current profound gap in understanding of human thymus development. Collectively, they demonstrate that the principal genetic mechanisms currently known to regulate thymus organogenesis in the mouse are conserved in humans, and that key cellular processes, including the developmental origin of the thymus in the third pharyngeal pouches, are also conserved. They establish that overt differentiation of cortical and medullary TECs occurs from mid-week 8 in the human thymic primordium and closely mirrors the progression seen in mouse in terms of phenotypic and functional marker expression, and that colonization of the human thymic primordium by endothelial and mesenchymal cells occurs from early and mid-week 8, respectively. Finally, they define the timing of onset of hematopoietic cell colonization of the human thymic primordium and reveal, in contrast to expectations from previous studies, that the first colonizing hematopoietic cells that express T lineage affiliated markers are CD45+CD7+CD34+ cells, which are present from early week 8. These findings are summarized in supplementary material Table S3.

**Conservation of mechanisms of thymus organogenesis between mouse and human**

The mouse is the principal model organism used to determine regulatory mechanisms operating to control thymus development and function, and therefore interrogation of its utility for predicting mechanisms that control these processes in the human is essential. The data presented above establish that the spatial and temporal expression pattern of all of the genes tested, which have been demonstrated by genetic analyses to have functional roles in thymus development, are conserved between mouse and human. They therefore validate the mouse as a model for understanding human thymus organogenesis. Of note is that although some variation in expression between mouse and human was observed in the genes analyzed herein, this did not affect expression in the thymus domain; for instance, PAX9 is expressed throughout the pharyngeal pouches in mouse but was detected only in pouches 3 and 4 in human. However, it is of interest that two genes, RHOX4 and PLET1, expression of which in mouse is restricted to the pharyngeal endoderm and the thymus domain of the third pharyngeal pouch, respectively, at this developmental stage, are not expressed in the human pharyngeal pouches or thymic rudiment, indicating that not all gene expression patterns are conserved. In this regard, we note that neither of these genes has been shown to be important for thymus organogenesis by functional studies and, indeed, both are currently of unknown function.

**Differentiation of the earliest progenitor cells in the human thymic primordium**

The data presented above demonstrate that overt differentiation of cortical and medullary TECs occurs at around mid-week 8. However, the thymic rudiment itself is present from early week 6 and therefore, our data further indicate that the week 6 and week 7 thymic primordia consist mainly of undifferentiated TECs. Thus, cell surface markers expressed by most TECs at these developmental stages will be useful for isolating and characterizing the differentiative potential of early progenitor TECs. Similarly, the expression patterns demonstrated here for CD205 and CLDN4 are
consistent with these markers identifying early medullary and cortical sub-lineage-restricted progenitors in the human, as well as mouse, thymus. Collectively, these data pave the way for prospective isolation and functional testing of defined human fetal TEC subpopulations, which will be essential if progress is to be made in defining conditions for in vitro propagation of human TEPCs or for generating these cells from embryonic stem cells for use in cell replacement therapies.

We note that FOXN1, a master regulator of TEC differentiation, is expressed in the human thymic primordia from mid-week 6. In the mouse, high-level Foxn1 expression is initiated at E11.5 and the onset of overt TEC differentiation closely follows this upregulation. The interval between upregulation of FOXN1 and the appearance of differentiating cTECs and mTECs is therefore much longer in human than in mouse. This probably reflects differences in cell cycle; however, it is also possible that other, as yet unknown mechanisms, limiting the time of onset of TEC differentiation are initiated or repressed with different kinetics to FOXN1 in the human.

Kinetics of initial mesenchymal and vascular endothelial cell entry into the human thymus

Our data indicate that entry of vascular endothelial progenitor cells into the human thymus occurs as early to mid-week 8, with migrating cells observed in the connective tissue between the carotid artery and the thymus from early to mid-week 7. They further show that, although formation of trabeculae is observed from mid-week 7, the time of entry of neural crest-derived mesenchymal cells into the epithelial regions of the thymus is mid-week 8. These processes are thus closely linked, as in the mouse. Notably, both events occur after initiation of high-level FOXN1 expression in the human thymus, consistent with the demonstration in mouse that thymic colonization with both mesenchymal and vascular endothelial cells is Foxn1 dependent. However, our data further suggest that entry of vascular endothelial progenitors into the epithelial region of the thymic primordium precedes its colonization with mesenchymal cells, in contrast to the widespread assumption that immigration of vascular endothelial cells follows that of neural crest-derived mesenchyme.

Identity of the first thymus-seeding hematopoietic cells

The data presented in Fig. 7 establish that hematopoietic cell colonization of the human thymus primordium occurs from early to mid-week 7. As in mouse, hematopoietic colonization of the thymus therefore closely follows upregulation of high level FOXN1 expression, consistent with FOXN1-regulation of CCL25 and DLL4, which are required for migration and T lineage commitment of thymus seeding progenitors, respectively (Liu et al., 2006; Hozumi et al., 2008; Koch et al., 2008).

The earliest human thymus seeding T lineage-restricted progenitors have previously been proposed to be CD45+CD34hiCD7+ cells, based on functional analysis of hematopoietic cells in the week 8–9 fetal thymus (Galay et al., 1993; Haynes and Heinly, 1995; Haddad et al., 2006). Our data indicate that the hematopoietic population of the early human fetal thymus is heterogeneous, and that all intrathymic hematopoietic cells are CD34hi until early to mid-week 9. Indeed, the first colonizing hematopoietic cells appearing at early week 7 are CD45+CD34+CD7+, and some CD45+ at least as early as early week 8 are Mac1+ macrophages. Intrathymic CD45+CD34hiCD7+ and CD45+CD34lo/–CD7+ cells are present from early week 8; this was surprising, because the earliest intrathymic T-lineage progenitors were expected to express CD34 (Galay et al., 1993; Haynes and Heinly, 1995; Haddad et al., 2006). Although we cannot exclude the possibility that these cells mature within the thymic primordium to generate CD45+CD34loCD7+ cells, we note that CD45+CD34loCD7+ cells first appear within the thymus at mid-week 9 coincident with establishment of blood vessels, and furthermore that cells of this phenotype are initially found closely associated with blood vessels. However, the thymus contains T-lineage progenitors at least as early as week 8, as evidenced by the robust development of thymocytes in fetal thymic organ cultures established at this developmental stage and the intrathymic expression of genes associated with early human thymocyte development. Thus, we favor the idea that the thymus-seeding population changes once the vasculature has been established as the primary route of entry, and comprises CD45+CD34loCD7+ cells only after this point.

CD45+CD34lo/–CD7+ cells have been described previously in the FL; these are NK cells, and also express CD56 (NCAM1 – Human Gene Nomenclature Database) (Phillips et al., 1992; Bárcena et al., 1993). Thus, the minor CD45+CD34lo/–CD7+ population present at week 8 might also be NK cells. The CD45+CD34loCD7hi population present in the pre-vascular thymus is, however, a novel intrathymic hematopoietic population that is probably T lineage committed based on its expression of CD7. In this regard, it is of interest that CD7hi cells were never detected in the connective tissue associated with the pre-vascular thymus or in the perithymic space, but were sometimes observed interspersed with thymic epithelial cells adjacent to the mesenchymal capsule, suggesting that in the pre-vascular thymus, upregulation of CD7 might occur upon contact with thymic epithelial cells. Of note is that the FL contains a population of CD45+CD34hiCD7– cells (Haddad et al., 2006) that might constitute precursors for this population. Based on these data, we therefore suggest that the pre-vascular thymus is initially seeded by CD45+CD34loCD7hi cells and that upregulation of CD7 in these cells occurs dynamically as part of the thymus-seeding process. We further suggest that this population is superseded by the previously described T lineage-restricted CD45+CD34loCD7+ population (Haddad et al., 2006) only following establishment of the thymic vasculature.

Acknowledgements

We thank Anne Sanderson, Isabel Morton, Joan Creiger and the staff of the Bruntsfield Suite of the Royal Infirmary of Edinburgh and the staff of the CASA clinic in Leiden for patient recruitment for provision of samples for these studies; Simon Monard for flow cytometry; and K. O’Neill (University of Edinburgh) and Sten Eirik Jacobsen (University of Oxford) for critical reading of the manuscript.

Funding

This work was funded by Leukaemia and Lymphoma Research [C.C.B., A.M.F.]; the Medical Research Council, UK [C.C.B., R.A.A.]; the Wellcome Trust [C.C.B., L.X.M., A.M.F.]; the EU FP7 integrated project EuroSystem [C.C.B., S.R.T.]; and the Netherlands Organisation for Scientific Research [Zon-MW Vidi grant #91710377 to T.C.]. Deposited in PMC for immediate release.

Competing interests statement

The authors declare no competing financial interests.

Supplementary material

Supplementary material available online at http://dev.biologists.org/lookup/suppl/doi:10.1242/dev.087320/-/DC1

References

Human thymus development


DEVELOPMENT


