A crucial role of WW45 in developing epithelial tissues in the mouse

Joo-Hyeon Lee1, Tae-Shin Kim1, Tae-Hong Yang1, Bon-Kyoun Koo2, Sang-Phil Oh1, Kwang-Pyo Lee1, Hyun-Jung Oh1, Sang-Hee Lee3, Young-Yun Kong2, Jin-Man Kim4 and Dae-Sik Lim1,*

1National Research Laboratory, Department of Biological Sciences, Korea Advanced Institute of Science and Technology, Daejeon, Korea; 2Division of Molecular and Life Sciences, Pohang University of Science and Technology, Kyungbuk, Korea; 3Division of Electron microscopic Research, Korea Basic Science Institute, Daejeon, Korea; 4Department of Pathology, College of Medicine, Chungnam National University, Daejeon, Korea

The role and molecular mechanisms of a new Hippo signalling pathway are not fully understood in mammals. Here, we generated mice that lack WW45 and revealed a crucial role for WW45 in cell-cycle exit and epithelial terminal differentiation. Many organs in the mutant mouse embryos displayed hyperplasia accompanied by defects in terminal differentiation of epithelial progenitor cells owing to impaired proliferation arrest rather than intrinsic acceleration of proliferation during differentiation. Importantly, the MST1 signalling pathway is specifically activated in differentiating epithelial cells. Moreover, WW45 is required for MST1 activation and translocation to the nucleus for subsequent LATS1/2 activation upon differentiation signal. LATS1/2 phosphorylates YAP, which, in turn, translocates from the nucleus into the cytoplasm, resulting in cell-cycle exit and terminal differentiation of epithelial progenitor cells. Collectively, these data provide compelling evidence that WW45 is a key mediator of MST1 signalling in the coordinate coupling of proliferation arrest with terminal differentiation for proper epithelial tissue development in mammals. The EMBO Journal (2008) 27, 1231–1242. doi:10.1038/emboj.2008.63; Published online 27 March 2008

Subject Categories: signal transduction; differentiation & death

Keywords: epithelial proliferation and differentiation; hWW45; MST1 signalling

Introduction

Homeostasis of regenerative epithelial tissues such as skin and intestine is maintained through a tightly balanced process of proliferation and terminal differentiation. During normal epithelial development, proliferating progenitor cells, often referred to as transiently amplifying cells, actively divide a limited number of times before they undergo cell-cycle exit and terminally differentiate into postmitotic cells (Blanpain et al., 2007). Cancer can develop as a result of inappropriate proliferation of progenitor cells accompanied by a partial or complete loss of differentiation (Reya et al., 2001). Therefore, understanding the signalling networks that control cell-cycle exit and terminal differentiation in epithelial tissues will provide insights into the mechanisms underlying tumorigenesis.

A new signalling network, known as the ‘Hippo pathway’ in Drosophila, seems to be a key developmental programme in controlling proliferation and apoptosis for proper organ development in Drosophila (Edgar, 2006; Harvey and Tapon, 2007; Pan, 2007; Saucedo and Edgar, 2007). The Ste-20 family kinase Hippo (Harvey et al., 2003; Jia et al., 2003; Pantalacci et al., 2003; Udan et al., 2003; Wu et al., 2003), WW adaptor protein Salvador (Kango-Singh et al., 2002; Tapon et al., 2002) and NDR kinase Warts (Justice et al., 1995; Xu et al., 1995) are the key components of the Hippo pathway that restrict cell proliferation and promote apoptosis in differentiating epithelial cells by regulating expression of cyclin E and Diap1. The Hippo kinase phosphoaylates and activates the Warts kinase, and this process is facilitated by the scaffolding protein Salvador or Mats (Wu et al., 2003; Wei et al., 2007). Warts, together with Mats, then phosphorylates and inhibits the transcription coactivator Yorkie (Huang et al., 2005; Lai et al., 2005). Expanded, Merlin and Fat, all of which localize to the plasma membrane, function upstream of the Hippo pathway (Bennett and Harvey, 2006; Cho et al., 2006; Hamaratoglu et al., 2006; Silva et al., 2006; Willecke et al., 2006). In flies, mutations of these factors lead to increased cell proliferation and decreased cell death.

The phenotypes of flies with mutations in the Hippo pathway can be rescued with their respective human counterparts (Tao et al., 1999; Wu et al., 2003; Huang et al., 2005; Lai et al., 2005), indicating that the Hippo pathway may have an analogous role in epithelial tissue development in mammals. Several reports on each mammalian component of the Hippo pathway have shown that the pathway is involved in cell death and cell-cycle regulation. MST1/2 kinases (Hippo homologues) were originally reported to be involved in apoptosis with caspase-3-mediated proteolytic activation (Lee et al., 1998). LATS1/2 (Warts homologues) have been implicated in the regulation of cell-cycle progression, apoptosis (Tao et al., 1999; Xia et al., 2002), mitotic exit and cytokinesis (McPherson et al., 2004; Yang et al., 2004). YAP (a Yorkie homologue) is involved in apoptosis by interacting with p73 (Matallanas et al., 2007). Although mutation of WW45 (a Salvador homologue) has been reported in several cancer cell lines (Tapon et al., 2002), little is known about the functional significance of WW45 in mammals. So far, only limited biochemical interactions have been reported, including the phosphorylation of LATS1/2 by MST1/2, the...
associations of WW45 with MST1/2 and LATS1/2, binding of LATS1 to MOB1 (a MAT3 homologue) and formation of a complex comprising RASSF1A, MST2, WW45 and LATS1 (Chan et al., 2005; Hergovich et al., 2006; Guo et al., 2007).

The Hippo pathway has also been implicated in mammalian tumorigenesis. Mice lacking LATS1 develop some types of tumour, and hWW45 and Mats are mutated in several cancer cell lines (St John et al., 1999; Tapon et al., 2002; Lai et al., 2005). NF2, the human orthologue of Merlin, is a tumour suppressor gene, mutations of which lead to neurofibromatosis (McClatchey and Giovannini, 2005). YAP is overexpressed in mammalian cancers and transgenic mice overexpressing YAP have an increased liver size and dysplasia with expanded undifferentiated progenitor cells in the intestine (Zender et al., 2006; Camargo et al., 2007; Dong et al., 2007). Of the Hippo pathway proteins, only LATS1-, LATS2-, NF2- and YAP-null mice have been generated; however, these mice are either early embryonic lethal or fail to recapitulate defects seen in the respective Drosophila mutants (McClatchey et al., 1997; St John et al., 1999; McPherson et al., 2004, Morin-kensicki et al., 2006). Therefore, compared with Drosophila, much less is known about the physiological function of the Hippo pathway in mammalian epithelial development. Furthermore, the molecular mechanisms by which this pathway is regulated during development are not fully understood in mammals.

In the present study, we generated mice lacking WW45 to examine the role of the Hippo pathway in mammals. Mutant embryos displayed unchecked proliferation and defects in terminal differentiation of epithelial cells. We also revealed the molecular mechanism by which MST1 signalling is spatiotemporally regulated to allow cell-cycle exit and activation of terminal differentiation in epithelial cells.

Results

Retarded growth and perinatal lethality of the WW45-deficient mice

To identify the role of WW45 in vivo, we generated WW45 mutant mice using embryonic stem (ES) cell technology. The targeted mutation replaced a 2.4-kb genomic region containing WW45 exon 2 with a puromycin cassette, leading to premature stop in WW45. After electroporation, the targeted clone was identified and transmitted through the germ line (Figure 1A). The absence of WW45 protein was confirmed by immunoblot assay (Figure 1B). Heterozygous mice were born healthy and fertile, and developed normally. However, only three dwarf homozygotes were found among 954 littersmates generated from their control littermates. WW45-/- embryos were found at embryonic days 17.5 (E17.5) and E18.5 (Supplementary Table 1). WW45-/- embryos up to E11.5 were morphologically indistinguishable from their control littermates. However, from about E13.5 onwards, WW45-/- embryos were slightly smaller than the controls, indicating a slower gain in body weight (Figure 1C). Despite growth retardation, we observed no consistent overt defects that would cause the embryonic lethality of mutants. This result prompted us to examine placentas from mutants and their littermates. WW45-null placentas displayed immature development with poor growth and vascularization of the labyrinth layers (Figure 1D). Defective intermingling of fetal and maternal vessels was confirmed by staining with anti-PECAM1 and anti-laminin (Supplementary Figure S1). Thus, the malfunctional labyrinth layer may affect the growth and viability of WW45-/- embryos.

Hyperplasia and immature differentiation of epithelial cells in the WW45-deficient embryos

Previous Drosophila studies have proposed important roles for SAV1 in the regulation of proliferation and apoptosis in epithelial tissues. Thus, we performed histological analyses of WW45-/- embryos at various embryonic stages. Interestingly, hyperproliferation of epithelial cells was clearly observed in the skin and intestine (Figures 2 and 3) and other organs (Supplementary Figure S2) of the WW45-/- embryos at E17.5.

We first characterized skin development in these WW45-/- mice. Dividing keratinocytes are normally restricted to the basal layer of wild-type epidermis, and as cells exit from the cell cycle, these keratinocytes move outwards and differentiate to form the spinous layers, the granular layers and the dead enucleated stratum corneum layers at the skin surface (Figure 2Aa). By contrast, the mutant epidermis had a more dense basal layer and the expanded suprabasal layers were less differentiated, with reduced enucleation and compaction of the developing granular cells (Figure 2Aa'). Development of hair follicles was rarely seen, and only small premature hair follicles were seen in null embryos at this stage. Co-staining for E-cadherin and Ki67 revealed that the mutant skin contained increased numbers of proliferative epithelial cells, compared with wild-type skin (Figure 2Ab, b'). Almost all the basal cells and several suprabasal cells expressed Ki67 in the mutant epidermis, whereas proliferation was restricted to the basal layer in the wild-type epidermis. TUNEL (terminal deoxynucleotidyl transferase biotin-dUTP nick-end labelling)-positive cells were also seen in wild-type epidermis but not in mutant epidermis (Figure 2Be, e'). Thus, increased proliferation in the suprabasal layer and repressed apoptosis of terminally differentiated keratinocytes contribute to hyperplasia in the epidermis of mutant embryos.

Epithelia of WW45-/- embryos were hyperproliferative but did not seem to undergo normal differentiation; therefore, we investigated whether epithelial differentiation was delayed and/or defective in mutant epithelia using a panel of antibodies against proteins that are expressed at defined stages of differentiation. Keratin 14 was normally expressed in one or two layers of basal cells in wild-type embryos, whereas it was strongly expressed in the multilayered basal cells in mutant embryos (Figure 2Ba, a'). Moreover, there were increased numbers of keratin-10-expressing cells in the suprabasal layers of the WW45-/- epidermis compared with wild-type embryos (Figure 2Bb'). Expression levels of loricrin and filaggrin, which are markers of late keratinocyte differentiation, were significantly downregulated in mutant epidermis, indicating defects in late differentiation (Figure 2Bc, d'). Skin-barrier development with X-gal staining further confirmed the absence of terminally differentiated layers (Supplementary Figure S3). Electron microscopy analysis clearly showed that the epidermis of WW45-null embryos was thicker than the wild-type epidermis. Moreover, the granular and cornified layers present dysmaturation in the mutant epidermis, with nucleated cells reaching the epidermal surface (Figure 2C). These data indicate that the
suprabasal mutant keratinocytes fail to stop proliferating and terminally differentiate.

We also examined intestinal development in mutant embryos. Wild-type intestinal epithelium consists of a monolayer of polarized epithelial cells organized into crypts. By contrast, the mutant epithelium was multilayered and displayed hypercellularity with pseudostratified and enlarged nuclei, perturbed differentiation with loss of goblet cells and increased numbers of mitotic cells (Figure 3Aa and Supplementary Figure S2a). Furthermore, Ki67 staining revealed extensive proliferation throughout the villus epithelium in the small intestine, whereas proliferative cells were restricted to the crypt bases in the control epithelium (Figure 2Ab, b). Staining with Alcian blue, a marker for goblet cells, revealed a complete absence of muco-secreting goblet cells in all mutant intestinal tracts (Figure 3Bc, c). Ultrastructural analysis also revealed poorly developed microvillus brush borders on the apical surfaces of the villous enterocytes, indicating defective enterocyte differentiation in the mutant epithelium of the small intestine (Figure 3C). Interestingly, these mutant cells had enlarged nuclei located close to the apical region, indicating a loss of apical–basal polarity.

In addition to immature differentiation of the mutant skin and intestine, immature differentiation was also detected in the lungs of mutant embryos (Lee and Lim, personal observation). Taken together, these results indicate that WW45 deficiency induces hyperplasia and immature differentiation in epithelial tissues.
WW45 regulates cell-cycle exit in epithelial progenitor cells during differentiation

We tested whether excessive proliferation of mutant cells was due to increased proliferation rates or failure of cell-cycle exit for terminal differentiation. First, we examined the cell-cycle duration in epithelial progenitor cells by analysing the proportion of cycling cells (Ki67⁺) in S-phase 1 h after injection of BrdU (Schmahl, 1983). Although the percentage of BrdU-labelled cells was increased in mutant embryos, the BrdU⁺ Ki67⁺/Ki67⁺ labelling index was approximately the same in wild-type and mutant embryos, indicating similar proliferation rates in wild-type and mutant embryos (Figure 4A). Second, we determined the frequency of cell-cycle re-entry by assessing the proportion of dividing cells (BrdU⁺) 24 h after BrdU injection (Chenn and Walsh, 2002). During the time interval between BrdU application and analysis, cells can leave (Ki67⁻) or re-enter the cell cycle (Ki67⁺). The mean ratio of BrdU⁺ Ki67⁺/BrdU⁺ cells was significantly increased by 49% in the mutant small intestine and by 58% in the mutant colon compared with wild-type controls (Figure 4B), indicating that WW45 promotes exit from the cell cycle in epithelial progenitors during embryonic development.

Failure of cell-cycle exit of WW45⁻/⁻ keratinocytes during in vitro differentiation

To further analyse the rates of proliferation and differentiation of epithelial cells, we isolated primary keratinocytes from the skin of embryos. Consistent with in vivo data (Figure 4A), WW45⁻/⁻ keratinocytes had normal cell-cycle distribution and the rate of proliferation was not significantly increased compared with control cells (Figure 4C and data not shown).
Again, this suggests that WW45 is unlikely to regulate the rate of proliferation. We then examined the ability of WW45 to regulate proliferation arrest and differentiation of developing epidermal cells by adding calcium, transforming growth factor (TGF)–β or LiCl, which have been shown to induce proliferation exit, and possibly terminal differentiation, of keratinocytes (Hennings et al., 1980; Shipley et al., 1986; Olmeda et al., 2003). With Ca²⁺, TGF-β or LiCl treatment, the wild-type cells showed efficient growth arrest and the numbers of BrdU-labelled cells were reduced. By contrast, mutant cells continued to proliferate and were BrdU-positive 24 h after Ca²⁺, TGF-β or LiCl treatment (Figure 4D and Supplementary Figure S5). Consistently, the expression of filaggrin was evident in wild-type but not in mutant keratinocytes after induction of differentiation (Figure 5A). These data indicate that increased proliferation in WW45−/− epithelial tissues results from impaired growth arrest of progenitor cells during differentiation rather than from an increased rate of proliferation.

**Activation of the MST signalling pathway during keratinocyte differentiation in vitro**

To further characterize the molecular mechanisms by which WW45 regulates cell-cycle exit during differentiation, we investigated the phosphorylation and localization of MST, LATS and YAP in keratinocytes during differentiation. Interestingly, autophosphorylation of MST1 was induced upon differentiation, as shown by phospho-MST1 immunoblotting, indicating MST1 activation (Figure 5A). By contrast, this autophosphorylation of MST1 was not detected in
Figure 4 Increased numbers of cycling cells in the developing WW45⁻/⁻ epithelium resulted from inefficient growth arrest. (A) The proportion of progenitor cells (Ki67⁺) labelled with BrdU after a 1 h pulse. There is no difference in cell-cycle length between wild-type and mutant embryos. (B) Analysis of immunoreactivity for Ki67 in BrdU-positive cells 24 h after BrdU injection. The fraction of cells re-entering the cell cycle (BrdU⁻ Ki67⁺) is significantly increased in mutant embryos. (C) Growth curves of primary keratinocytes isolated from wild-type and mutant epidermis show similar proliferation rates. (D) Induction of calcium-stimulated differentiation in primary keratinocytes. The level of BrdU incorporation in keratinocytes cultured in the absence or presence of Ca²⁺ for the times indicated is shown. The mutant keratinocytes show inefficient growth arrest under differentiation conditions. (A–D) Data represent triplicate independent experiments (± s.d.).

Figure 5 Activation of MST1 signalling pathway during differentiation. (A) Protein status of components of the MST1 pathway under differentiation conditions. Primary keratinocytes were incubated with or without Ca²⁺ for 24 h and analysed by western blot analysis with the indicated antibodies. Note the activation of MST1, as shown by pMST1 blotting, and the mobility shift of YAP and LATS1/2 in the control keratinocytes, but not in mutant keratinocytes, under differentiation conditions. Keratinocyte differentiation was confirmed by western blot analysis of filaggrin expression. (B) Wild-type and mutant primary keratinocytes cultured with or without Ca²⁺ for 24 h were analysed by immunoprecipitation with anti-MST1 and by western blot assays. Note that complex formation is only detected with Ca²⁺ treatment in the control samples. (C) Phosphorylation of YAP by activated LATS2 through MST1/WW45. WW45⁻/⁻ primary keratinocytes were co-transfected with the indicated plasmids and probed with the indicated antibodies. Note the mobility shift of YAP and LATS2 in the presence of MST1 and intact WW45.
mutant cells. Therefore, WW45 is required for MST1 activation after induction of differentiation with calcium treatment. In addition, the mobility patterns of LATS1/2 and YAP differ after induction of differentiation with calcium treatment (Figure 5A). Phosphorylation of LATS1/2 and YAP was detected in differentiated wild-type keratinocytes, but not in differentiated mutant keratinocytes. Thus, phosphorylation of YAP seems to be dependent on the MST1 signalling pathway, in particular on LATS1/2, during epithelial differentiation in mammals.

We then investigated whether the formation of components of the MST1 signalling pathway might be associated with and affect differentiation of keratinocytes. We first transfected WW45−/− keratinocytes with LATS1/2, MST1 and YAP with or without WW45, and then induced differentiation before immunoprecipitation. In mutant keratinocytes, LATS1 and LATS2 co-precipitated with YAP but not with MST1. By contrast, these proteins form a stable complex in the presence of WW45 (Supplementary Figure S6). Moreover, endogenous MST1 and LATS1/2 formed a complex in keratinocytes under differentiated conditions and in the presence of WW45 (Figure 5B). In addition, YAP was fully phosphorylated in keratinocytes complemented with wild-type WW45 and expressing LATS2 wild type, but not in cells with wild-type WW45 and expressing LATS2 kinase dead (KD) and MST1 or in cells complemented with WW45 lacking the SARAH domain, which is responsible for interaction with MST1 (Figure 5C). These results indicate that, in mammals, WW45 is required for MST1 activation and promotes LATS1/2 phosphorylation by recruiting MST1 into the complex, and that activated LATS1/2 then phosphorylates YAP. Surprisingly, activation of the MST1 signalling pathway in mammals seems to be specific for differentiation signals, at least for keratinocyte differentiation.

We next found that the serine 127 of YAP was phosphorylated by LATS1/2 in vitro and in vivo (Supplementary Figure S7). This result is consistent with the recent finding that serine 127 of YAP is the primary Hippo-responsive phosphorylation site (Dong et al., 2007). We further confirmed that this is a phosphorylation site of endogenous YAP with a phosphosine 127 antibody during keratinocyte differentiation (Supplementary Figure S7).

**Dynamic cellular localizations of MST1 and YAP during epithelial differentiation**

YAP associates with the Src family kinase at the plasma membrane, with 14-3-3 family proteins in the cytoplasm and with transcription factors in the nucleus, suggesting a dynamic localization of YAP in cells (Sudol, 1994; Yagi et al., 1999; Vassilev et al., 2001; Matallanas et al., 2007). However, the localization of other components of MST1 pathway has not been determined. Thus, we examined the subcellular localization of MST1 pathway components during epithelial cell differentiation by nuclear-cytoplasmic fractionation experiments. MST1 was mainly detected in the cytoplasmic fraction under undifferentiated conditions, but it was phosphorylated and mainly detected in the cytoplasm of differentiated wild-type cells. However, cytoplasmic localization and phosphorylation of YAP were compromised in mutant cells under differentiated conditions (Figure 6A). These results indicate that induction of keratinoctye differentiation triggers translocation of MST1 into the nucleus and then activates LATS1/2, which in turn phosphorylates YAP, and that WW45 is necessary for this. We further examined the ability of MST1/WW45/LATS2 to modulate the localization of YAP by ectopically overexpressing these proteins in mutant keratinocytes. Consistent with previous results, YAP was phosphorylated by LATS2 and this phosphorylated form was found in the cytoplasm, probably owing to nuclear translocation of MST1 by WW45 (Figure 6B). By contrast, these dynamic localizations of MST1 and YAP were not seen in cells expressing WW45 lacking the SARAH domain, which indicates that the WW45 SARAH domain is required for this process. Moreover, the YAP S127A mutant was mainly localized to the nucleus regardless of MST1 signalling activation, indicating that phosphorylation of serine 127 of YAP by LATS2 in the nucleus promotes translocation of YAP into the cytoplasm.

We also examined the distinct localization of MST1 and YAP by performing an immunostaining assay in differentiated keratinocytes as well as embryo tissue sections (Figure 6C and D). The specificity of MST1 antibody was assessed by immunoblot and immunostaining analyses with MST1-depleted cells (Supplementary Figure S8). Consistent with the fractionation results, we also observed dynamic translocation of endogenous YAP and MST1 during differentiation in the wild-type cells but such translocation was not apparent in the mutant cells during differentiation (Figure 6C). Moreover, although phosphorylated YAP was evidently found in the cytoplasm in differentiated wild-type cells, it was barely detectable in mutant cells. In wild-type tissue sections, we detected nuclear-localized YAP in proliferative basal cells, which were Ki67-positive, and cytoplasm-localized YAP in differentiated cells (Figure 6D and Supplementary Figure S9). This result is consistent with the recent finding that YAP is expressed and localized in the nuclei of the crypt compartment of the small intestine (Camargo et al., 2007). However, MST1 was detected in the cytoplasm of proliferating cells, but in the nucleus of differentiated cells in the wild-type skin and intestine. By contrast, many proliferating cells of mutant embryos showed cytoplasm-localized MST1 and nuclear-localized YAP (Figure 6D). In addition to changes in MST1 and YAP localizations, the cytoplasm of the differentiated regions in control embryos stained positive for phosphorylated YAP; but staining levels were significantly reduced in mutant embryos (Figure 6D). Therefore, consistent with in vitro epithelial differentiation results, the changes in the dynamic localization of MST1 and YAP are also likely to occur as epithelial cells undergo differentiation in vivo, and WW45 is likely to be a key protein in this process.

**The failure of proliferation arrest and differentiation of normal keratinocytes expressing YAP S127A**

We next assessed whether the phosphorylation of serine 127 of YAP is required for inducing keratinocyte growth arrest and differentiation. Thus, keratinocytes were infected with a retrovirus expressing YAP wild type, YAP S127A, which is a non-phospho form, or YAP S127D, which is a phospho-mimic
Figure 6 Dynamic localization of MST1/YAP during differentiation. (A–C) Subcellular localization of MST1 and YAP in response to differentiation stimuli. (A) Fractionation experiments were conducted with primary keratinocytes from wild-type or mutant embryos with or without Ca²⁺ for 24 h. Fractionated lysates were subjected to western blot analysis with the indicated antibodies. N, nuclear; C, cytoplasmic. (B) WW45-deficient keratinocytes were co-transfected with the indicated plasmids. After 24 h of transfection, cells were maintained in Ca²⁺ medium for a further 24 h before harvesting for fractionation experiments. Western blot analysis was performed with the indicated antibodies. Note that the major cytoplasm translocation of YAP and nuclear translocation of MST1 were detected only in the presence of intact WW45. (C) Primary keratinocytes were cultured with or without Ca²⁺ for 24 h and then subjected to immunostaining with anti-MST1, anti-YAP and anti-p-YAP. Note the phospho-dependent translocation of YAP and MST1 in Ca²⁺-induced differentiated control cells but not in mutant cells. (D) Immunoperoxidase staining for MST1, YAP and p-YAP in wild-type (a–f) and mutant (a’–f’) epithelium at E17.5. Note the intense nuclear detection of YAP and cytoplasmic detection of MST1 in most mutant epithelial cells, which is in contrast to the dynamic distributions according to differentiation stage in control epithelial cells. Serine 127 phospho-specific immunoreactivity was cytoplasm-specific in the differentiating zones in the control, whereas the signal was absent from the mutant cells. Counterstaining was performed with haematoxylin. (a–c, a’–c’) Skin; (d–f, d’–f’) small intestine. Scale bar: 10 μm (C) and 100 μm (D).
form. Calcium-induced differentiation signals suppressed proliferation of growing keratinocytes expressing YAP wild type or YAP S127D. By contrast, YAP S127A was mainly localized to the nucleus and its overexpression in wild-type cells failed to cause proliferation arrest and differentiation (Figure 7A–C). These results indicate that YAP S127A is localized to and acts in the nucleus to allow cells to proliferate even in the presence of differentiation signals. We also tested whether inactivation of YAP suppresses the differentiation defect in WW45-deficient keratinocytes.

To do this, we generated the dominant-negative form of YAP (DN), which dominantly inhibits the function of endogenous YAP protein (Zhao et al., 2007). Importantly, YAP DN was sufficient to rescue the differentiation of mutant cells (Figure 7D and E). Taken together, these results show that inactivation and subcellular targeting to the cytoplasm of phosphorylated YAP is required for cell-cycle exit and differentiation initiation.

**Figure 7** Effects of phosphorylation of YAP serine 127 on epithelial differentiation. (A) The percentage of proliferative cells labelled with BrdU under differentiation conditions. Levels of BrdU incorporation in primary keratinocytes were determined at the indicated times after retroviral infection with the indicated genes and incubation with or without Ca\(^{2+}\). WT, wild type; SA, S127A mutant; SD, S127D mutant. Note the failure of cell-cycle exit in response to Ca\(^{2+}\)-induced differentiation in YAP-SA-infected cells. (B) Failure of YAP-SA-infected cells to differentiate in response to differentiation stimuli. Lysates from (A) were subjected to western blot analysis with the indicated antibodies. Note the suppression of YAP mobility shift in the absence of processed forms of filaggrin in YAP-SA-infected cells. (C) Subcellular localization of YAP, YAP SA and YAP SD in response to differentiation stimuli. Fractionations from (A) were subjected to immunoblot analysis. Note the lack of cytoplasmic translocation of YAP SA under Ca\(^{2+}\)-induced differentiation conditions. (D) The percentage of BrdU-positive cells in mutant keratinocytes infected with the indicated genes under differentiation conditions. WT, wild type; DN, dominant-negative form. Note the suppression of cell-cycle re-entry in response to Ca\(^{2+}\) treatment in mutant keratinocytes infected with YAP-DN. (E) Lysates from (D) were subjected to western blot analysis with the indicated antibodies. Note the rescue of differentiation in response to Ca\(^{2+}\) treatment in mutant keratinocytes infected with YAP-DN. (F) Proposed model for the role of WW45 in developing epithelial tissues.
Discussion

This study provides novel insights into the role of WW45 in mammalian epithelial tissue development. WW45 is a key regulator of the MST1 signalling pathway, which promotes cell-cycle exit and terminal differentiation in developing epithelial tissues. Ablation of WW45 leads to hyperproliferation accompanied by immature differentiation in epithelial cells of the skin and intestine (Figures 2 and 3), and this phenotype results from re-entry of differentiating cells into the cell cycle rather than intrinsic acceleration of proliferation (Figure 4). This conclusion is strongly supported by our observations that WW45-null primary keratinocytes cannot be efficiently induced to exit the cell cycle in response to differentiation signals such as Ca^{2+}, TGF-β or LiCl, and that increased proliferation rates were detected from E15.5 onwards, which is the onset of terminal differentiation events in the epithelium (Figure 4 and Supplementary Figures S10 and S11). The Hippo pathway has been implicated in the restriction of proliferation and promotion of apoptosis in epithelial cells, but there is little evidence that this pathway is involved in terminal differentiation in Drosophila. Interestingly, the mouse MST1 signalling pathway seems to have a role in terminal differentiation in the developing epithelial tissues.

Previous studies with Drosophila have identified several genes and their interactions in the Hippo pathway (Edgar, 2006; Harvey and Tapon, 2007; Pan, 2007); however, the intracellular signalling of spatiotemporal regulation during epithelial differentiation remains unclear. This study determines the underlying mechanism by which components of the MST1 signalling pathway spatiotemporally regulate cell-cycle exit for epithelial differentiation in mammals. We show that as yet unknown differentiation signals specifically activate the MST1 kinase, which then dynamically localizes to the nucleus and activates LATS1/2, and that WW45 is required for this process. Activated LATS1/2 then phosphorylates serine 127 of YAP, and this phosphorylated YAP localizes to the cytoplasm (Figure 7F). Interestingly, transgenic mice overexpressing YAP develop liver cancers or OA P 

Recent studies in Drosophila have indicated that, together with cyclin E, Dia1 and bantam may be additional targets of Yorkie (Nolo et al., 2006; Thompson and Cohen, 2006). However, unlike in Drosophila, no direct target genes were identified in the MST1 pathway. Of interest, quantitative RT-PCR revealed that, although slightly increased, expression levels of cyclin E, cIAP, XIAP and survivin were not significantly upregulated in WW45-/- cells, indicating that other targets of YAP control developmental homeostasis in mammalian epithelial tissues (Supplementary Figure S12). Based on our observation that YAP DN is sufficient to rescue the cell-cycle exit and differentiation of WW45-/- cells, identification of the downstream targets or mediators of YAP will provide important insights into how YAP could control both cell-cycle exit and terminal differentiation.

We also found that WW45 deficiency led to disruption of contact inhibition of proliferation (data not shown), as is also seen with loss of LATS2 and NF2 (Lallemand et al., 2003; McPherson et al., 2004; Okada et al., 2005). WW45 is also likely to participate in the contact inhibition signalling pathway. Loss of contact inhibition might be associated with the hyperplasia observed in epithelial tissues of WW45-/- embryos. Recently, inactivation of YAP in the Hippo pathway has been shown to contribute to cell contact inhibition and tissue growth (Zhao et al., 2007). Thus, further studies should investigate whether defects in contact inhibition in WW45-/- cells are due to a failure of YAP inactivation.

Perturbations in MST1 signalling lead to inappropriate proliferation and expansion of cell compartments, which in turn lead to increased risks of cancer-associated mutations. Several results support the importance of the Hippo pathway in mammalian tumorigenesis. Loss of expression of LATS1/2, hWW45 and Mats has been reported in cancer cell lines (Tapon et al., 2002; Lai et al., 2005; Takahashi et al., 2005; Jiang et al., 2006). Mice with mutations in LATS1or NF2 develop tumours, and YAP transcription is increased in the mouse tumour model (St John et al., 1999; McClatchey and Giovannini, 2005; Zender et al., 2006). Moreover, YAP can transform immortalized mammary epithelial cells in vitro and transgenic mice overexpressing YAP develop liver cancers or intestinal dysplasia with loss of differentiated cell types (Overholtzer et al., 2006; Camargo et al., 2007; Dong et al., 2007). In addition to these reports, this study revealed that dysfunction of a single gene, WW45, affects most developing epithelial tissues and induces characteristics of a precancerous state: uncontrolled proliferation, partial loss of epithelial polarity and block of terminal differentiation. However, no such phenotypes have been reported in mice lacking LATS1, LATS2 and NF2. Multiple homologues of the fly Hippo pathway exist in mammals; hence, there may be functional redundancy of these proteins. However, WW45 is the only mammalian homologue of the Sav in the Hippo pathway, and its loss led to marked phenotypic changes in many epithelial tissues. Importantly, we found that, during a 14-month observation period, 22% of WW45 heterozygous mice developed some type of tumour including osteosarcoma and hepatoma (Lee and Lim, personal observation). Therefore, future studies should investigate how the heterozygous status of WW45 initiates tumorigenesis and the molecular characteristics of developed tumours in these mice. Finally, WW45-knockout embryo or WW45 heterozygous mice could be a suitable model for tumorigenesis studies of the MST1 pathway.
Materials and methods

Generation of WW45-null mice
WW45−/− mice were generated using standard ES cell (R1) homologous recombination and blastocyst injection techniques. A targeted ES cell clone was injected into C57BL/6 blastocysts to generate germline-transmitting chimeric mice. Mice and ES cells were genotyped by PCR assays with primers WW45-L (TGACCTGTCGACGCCCTTA), WW45-R (CGAATGCAGTCCTCATATTG) and pGK-3 (GCAGGACTAGTGACGTCGTAC).

Histological analysis
Embryos were fixed in 4% paraformaldehyde and embedded in paraffin. Sections (4 µm) were stained with H&E or subjected to immunohistochemical analysis. Immunohistochemical analysis was performed using standard protocols with antibodies against PECAM-1, laminin (Abcam), BrdU, E-cadherin, β-catenin (BD), Ki67 (Novocastra), anti-filaggrin, loricrin, K10, K14, K1 (Covance), iFABP (gift from Dr JI Gordon), chromogranin A (Immunostar), YAP (Cell Signalling, Santa Cruz), and p-YAP (Cell Signalling). Peroxidase levels were assessed using the EnVision™ + Dual Link System-HRP(DAB +) (DakoCytovention). Intestinal goblet cells were stained with Alcian blue to detect mucin. The TUNEL assay was performed using standard protocols.

Electron microscopy
Electron microscopy analysis of tissue samples was performed using a Tecnai G2 Spirit Twin high-voltage electron microscope (FEI Company, USA) and a JEM ARM 1300S high-voltage electron microscope (JEOL, Japan) were used.

Plasmid construction
Human cDNAs for WW45, MST1, LATS1/2 and YAP were cloned into pDK-Flag or pCMV-HA (HA: haemagglutinin), which had been modified from pcDNA3.1 or pcDNA3 (Invitrogen). Site-directed PCR mutagenesis was used to introduce the missense changes S127A and S127D into the YAP sequence. The S127A nuclear-localizing form with a deletion of the C-terminal TA domain of YAP was generated and used for YAP-DN mutant form.

Generation of antibodies
Rabbit polyclonal antibodies to MST1 were prepared with purified recombinant hexahistidine (His6)-tagged MST1(K59R) as the antigen and then affinity purified and used for immunostaining. Rabbit polyclonal antibodies to WW45 (Rb Ctrl#1) were generated by injecting rabbits with a keyhole-limpet-haemocyanin-conjugated peptide corresponding to the C-terminal amino acids of mouse WW45 (RKOQWQWAYQHCK). Specific antibodies were affinity purified with the appropriate antigens.

Primary keratinocyte cultures
Keratinocytes isolated from embryos were plated at 1.7×10^5 cells were cultured in 25 mL Tris–HCl (pH 7.4), 150 mM NaCl, 1 mM EDTA, 1 mM MgCl2, 0.2% Triton X-100, 0.3% NP-40, protease inhibitors and phosphatase inhibitors. Cell lysates were incubated for 2 h at 4°C with antibodies and then with protein A/G plus-agarose beads. Immunoprecipitates were subjected to western blot analysis. Separation of nuclear and cytoplasmic extracts was performed using NE-PER® Nuclear and Cytoplasmic Extraction Reagents (Pierce). Western blot analysis was performed using antibodies against YAP, p-YAP, MST1, p-MST1 (Cell Signalling), LATS1, LATS2 (Bethyl Laboratories Inc.), HA, filaggrin (Covance), laminin B (Santa Cruz), α-tubulin (Chemicon), Flag (Sigma) and WW45. For immunostaining analysis, keratinocytes were fixed and then exposed consecutively to primary and secondary antibodies. Slides were mounted with DAPI and imaged.

Supplementary data
Supplementary data are available at The EMBO Journal Online (http://www.embojournal.org).

Acknowledgements
We thank Dr H Saya for LATS1/2 cDNAs. This study was supported by grants from the NRI Program, the 21st Century Frontier Functional Human Genome Project, the Korea National Cancer Center Program and the Nuclear Research Grant.

References

© 2008 European Molecular Biology Organization

The EMBO Journal VOL 27 | NO 8 | 2008


Sudol M (1994) Yes-associated protein (YAP65) is a proline-rich phosphoprotein that binds to the SH3 domain of the Yes proto-oncogene product. Oncogene 9: 2145–2152


Tapon N, Harvey RF, Bell DW, Wahrer DC, Schiripo TA, Haber DA, Harlharian IK (2002) Salvador promotes both cell cycle exit and apoptosis in Drosophila and is mutated in human cancer cell lines. Cell 110: 467–478


