E4F1 connects the Bmi1-ARF-p53 pathway to epidermal stem cell-dependent skin homeostasis.

Julie Caramel, Matthieu Lacroix, Laurent Le Cam, Claude Sardet

To cite this version:
Julie Caramel, Matthieu Lacroix, Laurent Le Cam, Claude Sardet. E4F1 connects the Bmi1-ARF-p53 pathway to epidermal stem cell-dependent skin homeostasis. Cell Cycle, Taylor & Francis, 2011, 10 (6), pp.866-7. inserm-00610180

HAL Id: inserm-00610180
https://www.hal.inserm.fr/inserm-00610180
Submitted on 19 Mar 2012

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.
E4F1 connects the Bmi1-ARF-p53 pathway to epidermal stem cell-dependent skin homeostasis

Julie Caramel,1 Matthieu Lacroix,2* Laurent Le Cam2* and Claude Sardet1,*

1Institut de Génétique Moléculaire de Montpellier; UMR5535; CNRS; 2Institut de Recherche en Cancérologie de Montpellier; INSERM U896; Université Montpellier Sud de France; Montpellier, France

Skin homeostasis relies on epidermal stem cells (ESC) that are mobilized from their niche located in the basal layer of the interfollicular epithelium (IFE) and the bulge of hair follicles (HF) to fuel highly proliferative transit amplifying compartments (TAC). Differentiation programs then generate the spinous, granular and cornified layers of the IFE or the different lineages of the mature HF.1 ESC maintenance is orchestrated by a complex signaling network that remains incompletely characterized, including p63-, BMP-, TGFβ-, Wnt/β-catenin-, Rac1- and Notch-initiated signalling cascades.2 A recent work by Lacroix, Caramel et al. reveals that E4F1 and the Bmi1-Ink4a/Arf-p53 axis define a novel layer of regulation controlling ESC-dependent skin homeostasis (Fig. 1).2

E4F1 is an ubiquitously expressed multifunctional protein, originally identified as a transcription factor targeted by the adenoviral oncoprotein E1A. E4F1 also acts as an atypical ubiquitin E3-ligase for p53,3 and interacts with several components of the p53 pathway, including p14ARF and the polycomb member Bmi1.4

We recently generated E4F1 conditional knockout (KO) mice to circumvent the early embryonic developmental failure of E4F1 KO embryos3 and address E4F1 functions in adult tissues. E4F1 inactivation in adult skin first induces a transient hyperplasia of the epidermis associated with increased proliferation of keratinocytes with basal/TAC properties, followed by severe ulcerative lesions, broad disorganisation of the IFE, massive keratosis and ultimately, a permanent loss of cellularity of the epidermis and alopecia. E4F1 inactivation during embryonic development also resulted in similar progressive neo-natal skin defects, leading to dehydration and death of animals 3 to 4 days after birth.2

Strikingly, E4F1 KO skin defects result from cell autonomous perturbations in resident stem cells. Hence, various ESC markers were strongly downregulated in E4F1 KO epidermis [α6high/CD34+, K15+, BrdU Long-term Retaining Cells (LRC)] and E4F1 KO keratinocytes exhibited a dramatically impaired clonogenic potential ex vivo. These defects correlated with deregulated expression of the Ink4a/Arf locus in ESC clones and accordingly, E4F1 KO skin phenotypes were delayed in/mouse with Ink4a/Arf and E4F1 compound gene deficiencies. Inactivation of the Ink4a/Arf locus or ectopic expression of the Ink4a/Arf locus transcriptional repressor Bmi1 or depletion of its downstream target, p53, partly rescued the ex vivo clonogenic potential of E4F1 KO ESC.2 These results reveal a new regulatory network involved in ESC maintenance and proper skin homeostasis, implicating E4F1, Bmi1, the Ink4a/Arf locus and p53.

However, several questions remain unanswered. For instance, why are skin defects only developing after birth, although Cre-mediated recombination of E4F1 occurs much earlier during embryonic development? Among many possible scenarios, Bmi1 and its downstream effectors might impact on ESC maintenance only at post-natal stages, as already suggested for hematopoietic stem cells (HSC). Interestingly, E4F1 has been previously shown to interfere with this HSC program.3 However, in this case, the shRNA-mediated depletion of E4F1 rescues Bmi1 KO HSC exhaustion,4 suggesting that the role of the E4F1-Bmi1 axis in stem cells maintenance differs from one tissue to another. Moreover, while the E4F1-Bmi1 genetic interaction identified in HSC was claimed to be independent of p53,4 we found that p53 is clearly involved in E4F1 KO ESC phenotypes. The later observation raises queries about the exact function of the E4F1-Bmi1-Ink4a/Arf-p53 network in stem cells homeostasis.

In ESC, this pathway might play its usual “gate-keeper” function in response to genotoxic or developmental stresses, but it might also directly regulate normal programs of ESC self-renewal, as suggested in mammary gland progenitors5 where p53 impacts on asymmetric cell divisions.

Finally, given the multiple roles of E4F1, it will be necessary to evaluate the relative importance of E4F1-associated transcriptional and E3-ligase activities in ESC functions. Ongoing studies should clarify these points.

References

*Correspondence to: Laurent Le Cam and Claude Sardet; Email: Sardet@igmm.cnrs.fr and laurent.lecam@inserm.fr
Submitted: 01/20/11; Accepted: 01/28/11
DOI: 10.4161/cc.10.6.14974
Figure 1. E4F1, through its connection with the Bmi1-ARF-p53 axis, regulates epidermal stem cell (ESC) maintenance. Constant renewal of the Interfollicular Epithelium (IFE) and of Hair Follicles (HF) relies on the recruitment of epidermal stem cells (ESC) located in the basal layer of IFE and in the bulge region of HF, respectively. ESC fuel the highly proliferative transit amplifying compartments (TAC) in the basal layer of IFE and in the bulb of HF. TAC cells then embark on differentiation programs to generate the spinous, granular and cornified layers in IFE or the different lineages of mature HF. E4F1 inactivation in the entire skin or in the basal compartment of the epidermis induces skin homeostasis defects that result from cell autonomous alterations in ESC maintenance. These ESC defects are partially restored upon inactivation of the Bmi-Arf-Mdm2-p53 pathway, several components (p53, Bmi1 and Arf) of which have been described to physically and functionally interact with E4F1.