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Erratum
KLF5 mediates the hyper-proliferative phenotype of the intestinal epithelium in mice with intestine-specific endogenous K-Ras^{G12D} expression: Am J Cancer Res. 2018; 8(4): 723-731

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In the article AJCR0075555, errors have been found in the published versions of **Figures 3** and **5**. There are duplicate images presented in **Figures 3Ba**, **3Cb**, and **5N**. To ensure integrity, we redid staining for KLF5 and pMEK on slides from *LSL-KRas* and *Villin-Cre/LSL-KRas* mice. We have accordingly corrected and replaced images in **Figure 3Ba**, **3Be**, **3Cb**, and **3Ch**, and kept **Figure 5N** unchanged. The representative images included in the revised version of **Figure 3** do not affect the interpretation of the results. The errors do not bear on the interpretation of the results, nor do they influence the conclusion of the manuscript. We sincerely apologize for this oversight and any confusion it may have caused. The corrected **Figure 3** is shown below.

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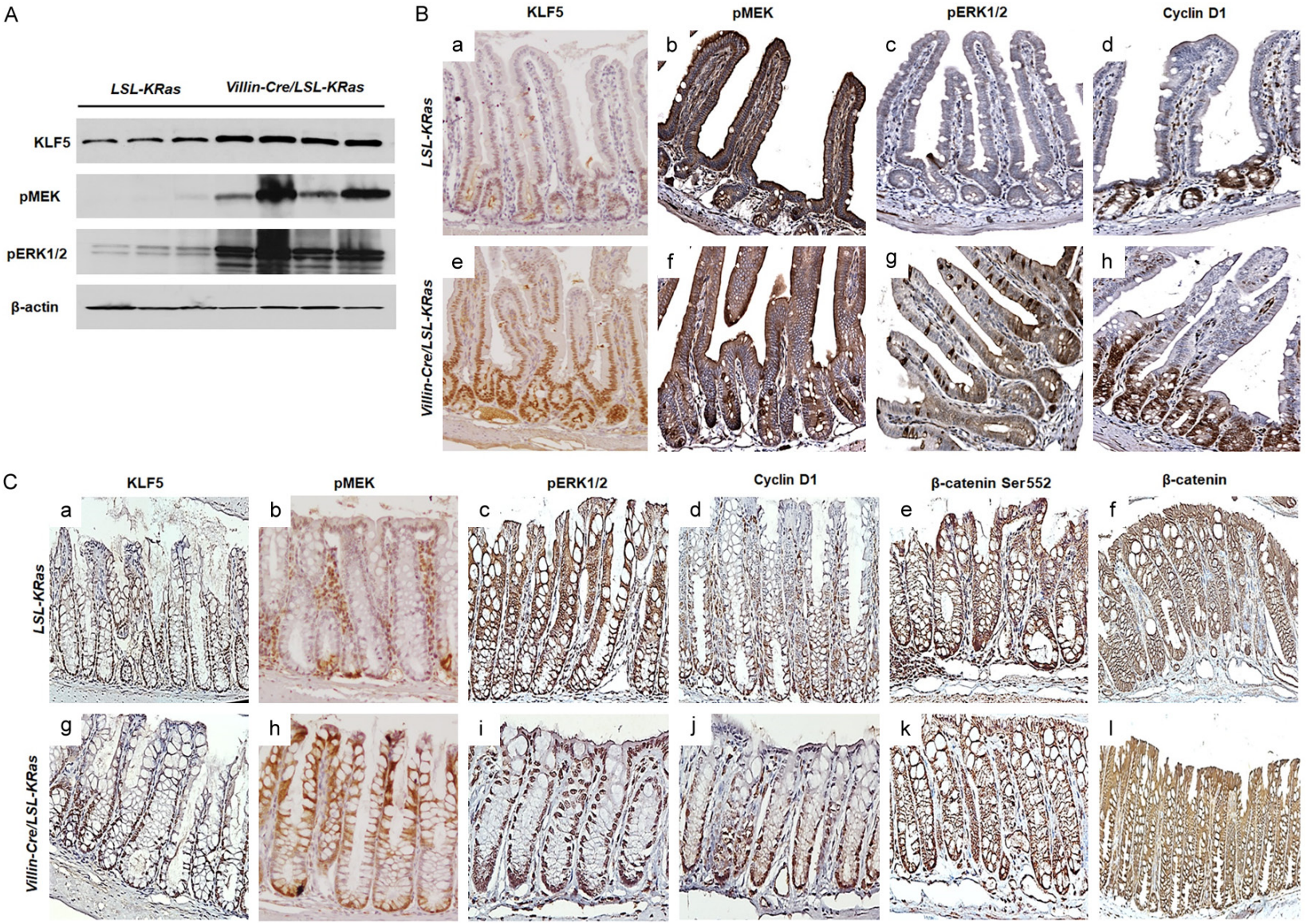

Figure 3. Upregulation of downstream targets of ERK pathway. A. Western Blot analysis of protein extract from normal appearing mucosa obtained from small intestines tissues from *LSL-K-Ras*^{G12D} and *Villin-Cre/LSL-K-Ras*^{G12D} mice. 10 ug of total protein extracts from *LSL-K-Ras*^{G12D} and *Villin-Cre/LSL-K-Ras*^{G12D} were analyzed by Western blotting using antibody against KLF5, pMEK, pERK1/2, Cyclin D1 and β-actin (as a loading control). The images are shown separately as they were obtained from different gels or different parts of the same gel. B. Small intestinal tissues from *LSL-K-Ras*^{G12D} and *Villin-Cre/LSL-K-Ras*^{G12D} mice were stained with antibody against KLF5 (a, e), pMEK (b, f), pERK1/2 (c, g) and Cyclin D1 (d, h). C. Colonic tissues from *LSL-K-Ras*^{G12D} and *Villin-Cre/LSL-K-Ras*^{G12D} mice were stained with antibody against KLF5 (a, g), pMEK (b, h), pERK1/2 (c, i), Cyclin D1 (d, j), β-catenin Ser552 (e, k), and whole β-catenin (f, l).