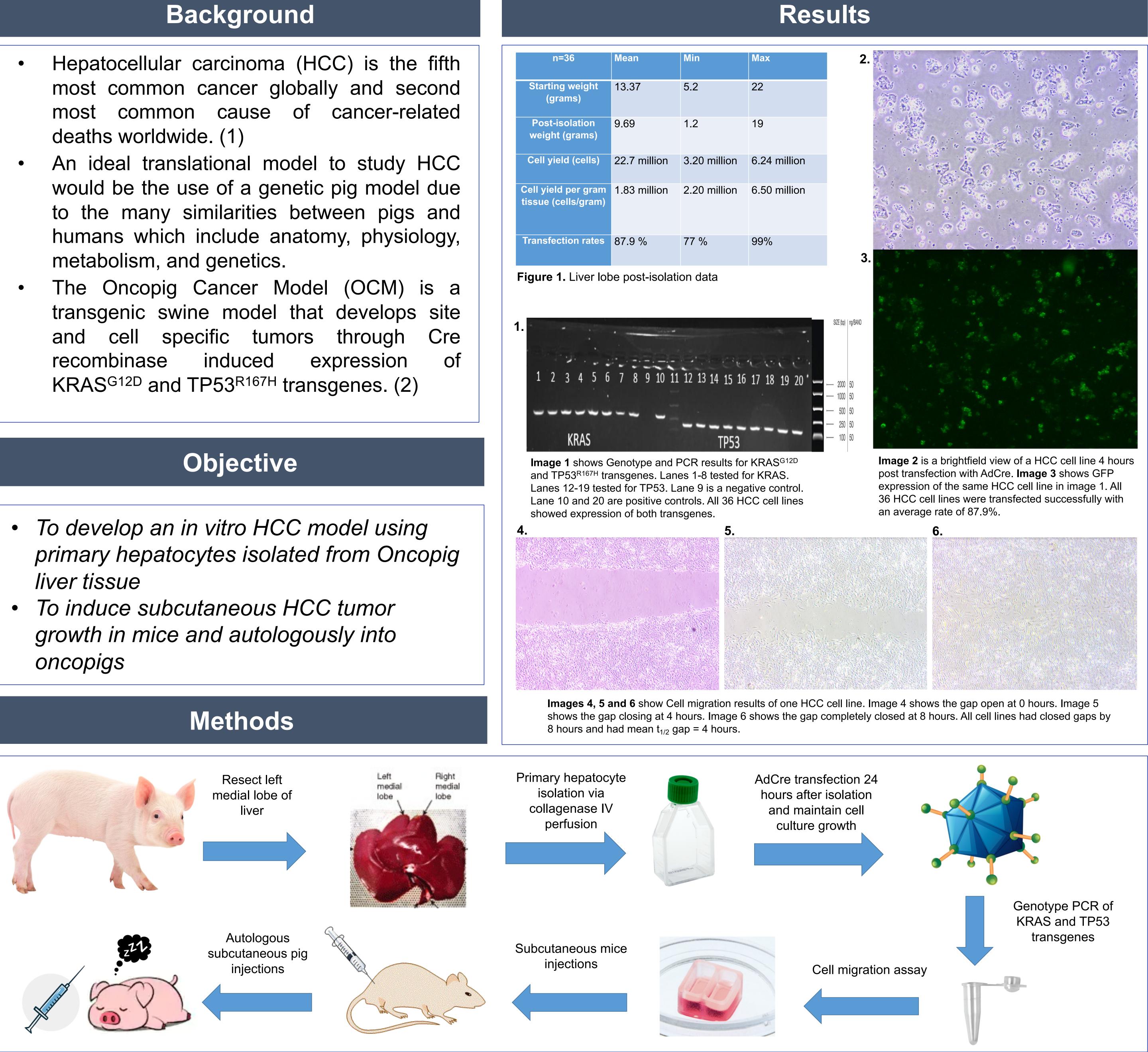


- most deaths worldwide. (1)
- metabolism, and genetics.
- specific tumors through cell and recombinase induced expression KRAS^{G12D} and TP53^{R167H} transgenes. (2)

- To develop an in vitro HCC model using liver tissue
- oncopigs



Development, Maintenance, and Characterization of Porcine Hepatocellular Carcinoma Cell Lines

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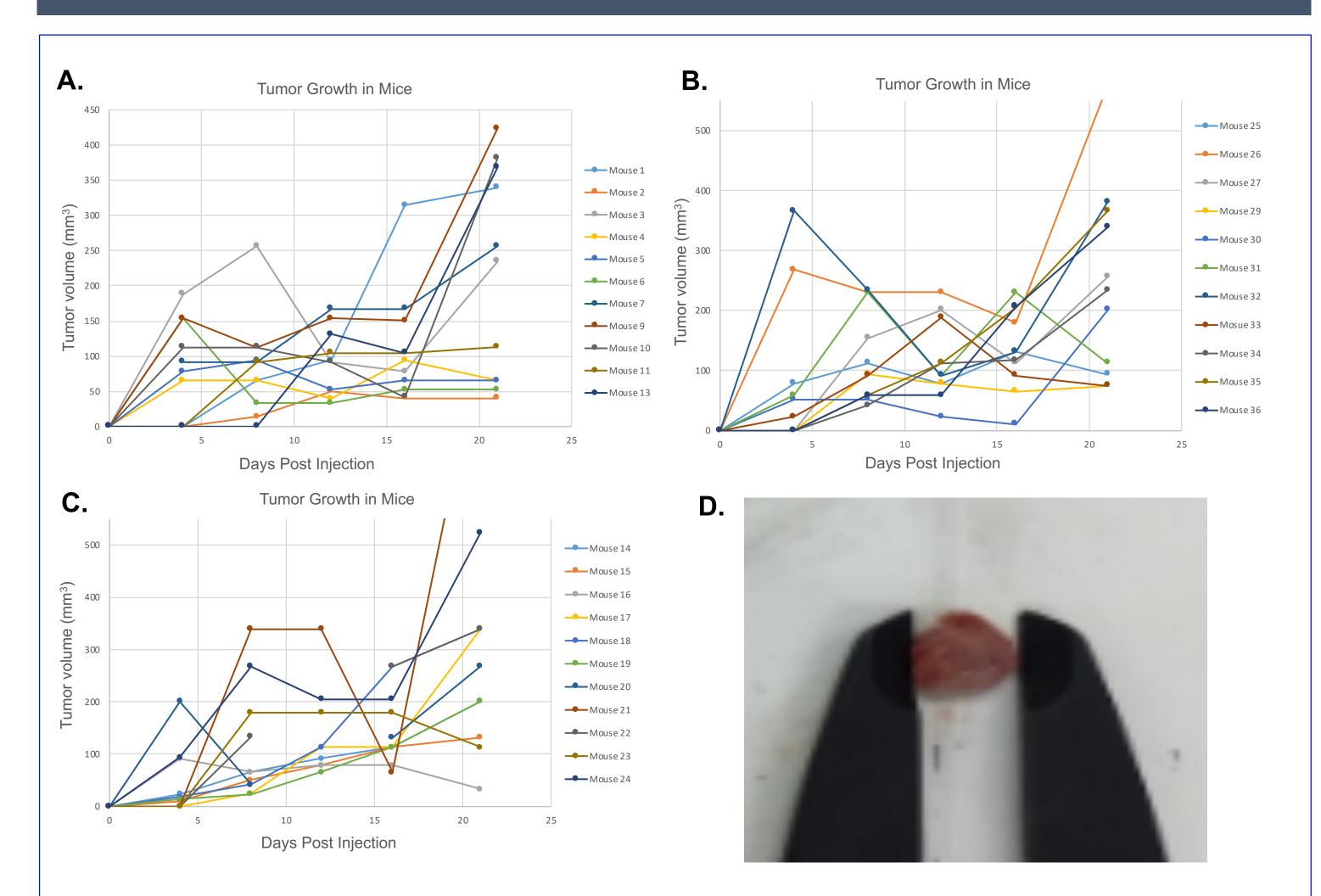


Figure 2. Graphs A, B and C show tumor volume growth in mice post injection (n= 34). Image D shows one collected tumor from a mouse. SCID or NSG mice received 2 subcutaneous injections in the abdominal region. Each injection contained 5 million porcine HCC cells per injection. Tumor growth was measured over the course of 21 days. Each mouse was euthanized on the 21st day and tumors were collected. A total of 59 tumors were observed and collected from 33 mice (86.8% tumor growth success). Mean volume for the tumors was 285.3 mm³ ranging from 18.84 mm³ to 1766.3 mm³.



Conclusions & Future Work

- and characterized by in vitro assays.
- oncopigs.
- techniques for HCC.
- (2018). 29(8), 1194-1202

Results

Image 7. Oncopigs (n=29) received a total of 6 subcutaneous injections autologously in the thoracic region. Each injection contained 10 million porcine HCC cells per injection. A total of 80 tumors were observed among 20 oncopigs (46% tumor growth success).

• A genetic swine model for HCC can be maintained by cell culturing

• Subcutaneous HCC tumor growth can be induced in mice and

• Future work involves developing intrahepatic tumors in oncopigs in order to further improve translational research in treatment

References

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