

Abstract

In vitro skin models have numerous applications in the field of pharmacology, toxicology and in understanding various disease mechanisms. Extracellular matrices from porcine gall bladder, small intestine and urinary bladder have been shown to have excellent scaffold properties for tissue engineering applications due to the presence of cytokines and growth factors that provide the biological signals for wound healing^{2,3}. HaCaT cells, having stable genetic character, can be effortlessly exploited for in vitro skin studies as they are easier to grow and subculture when compared to primarily isolated keratinocytes. Here, an in-vitro skin model was constructed using HaCaT cells seeded on three prototypes of porcine ECM scaffolds for skin tissue engineering and regenerative medicine applications⁴ and it was characterized structurally and functionally.

Methodology

Three prototypes of tissue engineering scaffolds were prepared from porcine gall bladder (cholecyst), small intestine and urinary bladder using previously published protocol⁴ and named as cholecyst, jejuna and urinary-bladder derived scaffolds (or C34 C34 der (or 9.224 25c2F 9.27v(de)23(r)-8()-654()17(or)-8(9.224 25c2F 9.27v(de)23(r)-8()-65