cultured using a non-serum media formulation. In many cases, the population of cells is cultured using a mushroom-based media formulation.

[038] Disclosed herein, in some examples, are cultured food products for human consumption, comprising the cultured tissue produced according to the methods of any one of the foregoing methods. Sometimes, the cultured food product comprises packaging having a label indicating the cultured tissue was produced in a pathogen-free environment, a toxin-free environment, without force-feeding an animal, or any combination thereof. In certain instances, the cultured tissue is processed into a plurality of slices and packaged to form the cultured food product.

[038a] In one aspect, disclosed herein is a method of producing cultured tissue for human consumption, the method comprising:

a) obtaining a population of self-renewing cells derived from salmon;
b) culturing the population of self-renewing cells derived from salmon for suspension culture in a bioreactor;
c) inducing differentiation in the population of self-renewing cells into myocytes, adipocytes, or a combination thereof to form cultured tissue; and

processing the cultured tissue for human consumption.

[038b] In one aspect, disclosed herein is a sushi grade salmon composition comprising the cultured tissue as described herein.

BRIEF DESCRIPTION OF THE DRAWINGS

[039] A better understanding of the features and advantages of the present invention will be obtained by reference to the following detailed description that sets forth illustrative embodiments, in which the principles of the invention are utilized, and the accompanying drawings of which:

[040] **FIG. 1** shows a flow chart for a process of producing cultured cells having lipid accumulation for human consumption.

CLAIMS

WHAT IS CLAIMED IS:

1. A method of producing cultured tissue for human consumption, the method comprising:
    a) obtaining a population of self-renewing cells derived from salmon;
    b) culturing the population of self-renewing cells derived from salmon for suspension culture in a bioreactor;
    c) inducing differentiation in the population of self-renewing cells into myocytes, adipocytes, or a combination thereof to form cultured tissue; and
    d) processing the cultured tissue for human consumption.
2. The method of claim 1, wherein culturing comprises seeding the population of self-renewing cells on 3-dimensional scaffolds.
3. The method of claim 2, wherein the 3-dimensional scaffolds are conjugated to at least one factor promoting cell growth, adhesion, differentiation, or a combination thereof, wherein the scaffolds comprise at least one of hydrogel, chitosan, polyethylene terephthalate, collagen, elastin, heparan sulfate, chondroitin sulfate, keratan sulfate, hyaluronic acid, laminin, fibronectin, cellulose, hemicellulose, pectin, lignin, alginate, glucomannan, polycaprolactone (PCL), textured vegetable protein (TVP), textured soy protein (TSP), and acrylates.
4. The method of claim 1, wherein at least one cell has been modified to incorporate:
    a) a first genetic construct comprising an open reading frame (ORF) of at least one pluripotency gene; and
    b) a second genetic construct comprising an open reading frame (ORF) of a regulatory factor configured to inactivate the at least one pluripotency gene.
5. The method of claim 4, wherein the regulatory factor is a recombinase, and the open reading frame (ORF) of at least one pluripotency gene is flanked by recombination sequences recognized by the recombinase such that expression of the recombinase catalyzes excision of the open reading frame (ORF) of at least one pluripotency gene.

6. The method of claim 4, wherein the second genetic construct comprises at least one myogenic factor selected from Myogenin (MyoG), Myogenic Differentiation 1 (MyoD), Myogenic Factor 6 (MRF4), and Myogenic Factor 5 (MYF5); or wherein the second genetic construct comprises at least one adipogenic factor selected from Fatty Acid Binding Protein 4 (FABP4), Insulin-Responsive Glucose Transporter Type 4 (GLUT4), Adiponectin, C1Q And Collagen Domain Containing (ADIPOQ), 1-Acylglycerol-3-Phosphate O-Acyltransferase 2 (AGPAT2), Perilipin 1 (PLIN1), Leptin (LEP), and Lipoprotein Lipase (LPL).
7. The method of claim 4, wherein the second genetic construct further comprises:
    a) an open reading frame (ORF) of at least one differentiation gene; and
    b) an inducible promoter controlling expression of:
        i. the open reading frame (ORF) of the at least one differentiation gene; and
        ii. the open reading frame (ORF) of the regulatory factor.
8. The method of claim 7, wherein inducing differentiation comprises exposing the at least one cell to an induction agent to induce expression of the ORF of the at least one differentiation gene and the ORF of the regulatory factor, and further comprising removing the induction agent after the population of self-renewing cells has been treated with the induction agent and before being processed for human consumption in d).
9. The method of claim 1, wherein inducing differentiation comprises at least one of generating myotubes within the population of self-renewing cells or generating adipocytes within the population of self-renewing cells.
10. The method of claim 1, wherein the population of self-renewing cells comprises multipotent cells that are induced to differentiate into myocytes and adipocytes during c) or a first subpopulation of myosatellite cells and a second subpopulation of pre-adipocytes.
11. The method of claim 1, wherein the cultured tissue comprises fish muscle tissue.
12. The method of claim 1, wherein the population of cells is cultured using a non-serum media formulation.
13. A sushi grade salmon composition comprising the cultured tissue of claim 1.