WHAT IS CLAIMED IS:

1. A pharmaceutical composition comprising:

   (a) a first component comprising a combination of (i) mycophenolic acid or a pharmaceutically acceptable salt, derivative or analog thereof, and (ii) a corticosteroid or a pharmaceutically acceptable salt, derivative or analogs thereof, wherein the corticosteroid is present at a concentration of about 0.005 % w/w to 0.05 % w/w; and

   (b) a carrier selected from the group consisting of de-ionized water and a balanced salt solution.

2. The pharmaceutical composition of claim 1, wherein the pharmaceutically acceptable salt of mycophenolic acid is mycophenolate sodium or mycophenolate mofetil.

3. The pharmaceutical composition of claim 1, wherein the corticosteroid is betamethasone.

4. The pharmaceutical composition of claim 3, wherein the salt, derivative or analog of betamethasone is selected from the group consisting of betamethasone acetate, betamethasone sodium phosphate, betamethasone valerate, and betamethasone dipropionate.

5. The pharmaceutical composition of claim 1, further comprising at least one second component selected from the group consisting of glycerol, polyvinylpyrrolidone, sorbitol, polyethylene glycol, hydroxypropyl methylcellulose, carboxymethylcellulose, and polyvinyl acetate.

6. The pharmaceutical composition of claim 5, wherein the concentration of the second component is between about 0.1 % w/w and about 5.0 % w/w.

7. The pharmaceutical composition of claim 5, further comprising at least one third component selected from the group consisting of a glycosaminoglycan, chondroitin, chondroitin sulfate, dermatan sulfate, dermatin sulfate, heparin sulfate, heparan sulfate, keratin sulfate, keratan sulfate, and hyaluronic acid.

8. The pharmaceutical composition of claim 7, wherein the concentration of the third component is between about 0.1 % w/w and about 5.0 % w/w.

9. The pharmaceutical composition of claim 7, further comprising at least one fourth component selected from the group consisting of dextran, dextran sulfate, sodium chloride, dextrose, and sucrose.

10. The pharmaceutical composition of claim 9, wherein the concentration of the fourth component is between about 0.1 % w/w and about 5.0 % w/w.

11. The pharmaceutical composition of claim 1, further comprising at least one additional compound selected from the group consisting of tacrolimus, lifitegrast, cyclosporine, a corticosteroid other than betamethasone, pharmaceutically acceptable salts, derivatives or analogs thereof, albumin, plasma, platelet-rich plasma, and serum.

12. The pharmaceutical composition of claim 11, wherein the corticosteroid other than betamethasone is selected from the group consisting of loteprednol, triamcinolone, triamcinolone acetonide, triamcinolone diacetate, triamcinolone benetonide, triamcinolone furetonide, triamcinolone hexacetonide, dexamethasone, fluorometholone, fluocinolone acetonide, prednisone, prednisolone, methylprednisone, corticol, cortisone, fluorocortisone, deoxycorticosterone acetate, aldosterone, and budesonide.

13. The pharmaceutical composition of claim 1, further comprising at least one antioxidant selected from the group consisting of ascorbic acid, erythorbic acid, sodium ascorbate, thioglycerol, cysteine, acetylcysteine, cystine, dithioerythreitol, dithiothreitol, glutathione, tocopherols, butylated hydroxyanisol, butylated hydroxytoluene, sodium sulfate, sodium bisulfite, acetone sodium bisulfite, sodium metabisulfite, sodium sulfite, sodium formaldehyde sulfoxylate, sodium thiosulfate, and nordihydroguaiaretic acid.

14. The pharmaceutical composition of claim 1, wherein the mycophenolic acid is present at a concentration of about 0.05 % w/w to about 1.0 % w/w.

15. The pharmaceutical composition of claim 14, wherein the first component includes mycophenolic acid at 0.3 % w/w and betamethasone at 0.03 % w/w.

16. The pharmaceutical composition of claim 1, wherein the concentration of the first component is between about 0.001 % w/w and about 75.0 % w/w.

17. The pharmaceutical composition of claim 1, further comprising at least one excipient or surfactant selected from the group consisting of a non-ionic polyoxyethlene-polyoxypropylene block copolymer, methylcellulose, a hydroxypropyl methylcellulose polymer, a polycarbophil polymer, edetate disodium dihydrate (EDTA), polysorbate-80, sodium thiosulfate, sodium thiosulfate pentahydrate, sodium phosphate, sodium phosphate dibasic anhydrous, and sodium phosphate monobasic anhydrous.

18. The pharmaceutical composition of claim 17, wherein the concentration of the excipient or surfactant is between about 0.1 % w/w and 4.0 % w/w.

19. The pharmaceutical composition of claim 1, wherein the corticosteroid is cyclosporine and wherein the composition further comprises:

(a) at least one second component selected from the group consisting of glycerol, polyvinylpyrrolidone, sorbitol, polyethylene glycol, hydroxypropyl methylcellulose, carboxymethylcellulose, and polyvinyl acetate;

(b) at least one third component selected from the group consisting of a glycosaminoglycan, chondroitin, chondroitin sulfate, dermatan sulfate, dermatin sulfate, heparin sulfate, heparan sulfate, keratin sulfate, keratan sulfate, and hyaluronic acid; and

(c) at least one excipient or surfactant selected from the group consisting of a non-ionic polyoxyethlene-polyoxypropylene block copolymer, methylcellulose, a hydroxypropyl methylcellulose polymer, a polycarbophil polymer, edetate disodium dihydrate (EDTA), polysorbate-80, sodium thiosulfate, sodium thiosulfate pentahydrate, sodium phosphate, sodium phosphate dibasic anhydrous, and sodium phosphate monobasic anhydrous.

20. The pharmaceutical composition of claim 2, wherein the pharmaceutically acceptable salt of mycophenolic acid is mycophenolate sodium and the corticosteroid is loteprednol etabonate, and wherein the composition further comprises:

(a) at least one second component selected from the group consisting of glycerol, polyvinyl pyrrolidone, sorbitol, polyethylene glycol, hydroxypropyl methylcellulose, carboxymethylcellulose, and polyvinyl acetate;

(b) at least one third component selected from the group consisting of a glycosaminoglycan, chondroitin, chondroitin sulfate, dermatan sulfate, dermatin sulfate, heparin sulfate, heparan sulfate, keratin sulfate, keratan sulfate, and hyaluronic acid; and

(c) at least one excipient or surfactant selected from the group consisting of a non-ionic polyoxyethlene-polyoxypropylene block copolymer, methylcellulose, a hydroxypropyl methylcellulose polymer, a polycarbophil polymer, edetate disodium dihydrate (EDTA), polysorbate-80, sodium thiosulfate, sodium thiosulfate pentahydrate, sodium phosphate, sodium phosphate dibasic anhydrous, and sodium phosphate monobasic anhydrous.

21. The pharmaceutical composition of claim 2, wherein the pharmaceutically acceptable salt of mycophenolic acid is mycophenolate sodium and the derivate or analog of the corticosteroid is betamethasone sodium phosphate, and wherein the composition further comprises:

(a) at least one second component selected from the group consisting of glycerol, polyvinylpyrrolidone, sorbitol, polyethylene glycol, hydroxypropyl methylcellulose, carboxymethylcellulose, and polyvinyl acetate;

(b) at least one third component selected from the group consisting of a glycosaminoglycan, chondroitin, chondroitin sulfate, dermatan sulfate, dermatin sulfate, heparin sulfate, heparan sulfate, keratin sulfate, keratan sulfate, and hyaluronic acid;

(c) at least one fourth component selected from the group consisting of dextran, dextran sulfate, sodium chloride, dextrose, and sucrose; and

(d) at least one excipient or surfactant selected from the group consisting of a non-ionic polyoxyethlene-polyoxypropylene block copolymer, methylcellulose, a hydroxypropyl methylcellulose polymer, a polycarbophil polymer, edetate

disodium dihydrate (EDTA), polysorbate-80, sodium thiosulfate, sodium thiosulfate pentahydrate, sodium phosphate, sodium phosphate dibasic anhydrous, and sodium phosphate monobasic anhydrous.

22. The pharmaceutical composition of claim 2, wherein the pharmaceutically acceptable salt of mycophenolic acid is mycophenolate sodium and the derivate or analog of the corticosteroid is tacrolimus monohydrate, and wherein the composition further comprises:

(a) at least one second component selected from the group consisting of glycerol, polyvinylpyrrolidone, sorbitol, polyethylene glycol, hydroxypropyl methylcellulose, carboxymethylcellulose, and polyvinyl acetate;

(b) at least one third component selected from the group consisting of a glycosaminoglycan, chondroitin, chondroitin sulfate, dermatan sulfate, dermatin sulfate, heparin sulfate, heparan sulfate, keratin sulfate, keratan sulfate, and hyaluronic acid;

(c) at least one fourth component selected from the group consisting of dextran, dextran sulfate, sodium chloride, dextrose, and sucrose;

(d) at least one antioxidant selected from the group consisting of ascorbic acid, erythorbic acid, sodium ascorbate, thioglycerol, cysteine, acetylcysteine, cystine, dithioerythreitol, dithiothreitol, glutathione, tocopherols, butylated hydroxyanisol, butylated hydroxytoluene, sodium sulfate, sodium bisulfite, acetone sodium bisulfite, sodium metabisulfite, sodium sulfite, sodium formaldehyde sulfoxylate, sodium thiosulfate, and nordihydroguaiaretic acid; and

(e) at least one excipient or surfactant selected from the group consisting of a non-ionic polyoxyethlene-polyoxypropylene block copolymer, methylcellulose, a hydroxypropyl methylcellulose polymer, a polycarbophil polymer, edetate disodium dihydrate (EDTA), polysorbate-80, sodium thiosulfate, sodium thiosulfate pentahydrate, sodium phosphate, sodium phosphate dibasic anhydrous, and sodium phosphate monobasic anhydrous.