

## **Rule 40-5-1-.02 Adoption by Reference**

Hereinafter, the following is adopted by reference, and therefore all applicable provisions become part of this chapter:

1. Federal Food, Drug and Cosmetic Act, Title 21 United States Code Parts 321 (Only 21 USC §§ 321(f), 321(g), 321(k), 321(m), 321(s), 321(v), and 321(w)), 331 (Only 21 USC §§ 331(a)-(f) and 331(k)), 333, 334, 341, 342 (Only 21 USC §§ 342(a)-(c)), 343 (Only 21 USC §§ 343(a)-(n)), 344, 346, 346a, 348, 351, 354, 360b, 371, and 374.
2. The Code of Federal Regulations, Title 21 CFR Parts 1 (Only 21 CFR §§ 1.20-.24), 7 (Only 21 CFR §§ 7.1-7.13 and 21 CFR §§ 7.40-7.59), 70 (Only 21 CFR §§ 70.20-70.25), 73 (Only 21 CFR §§ 73.1-73.615), 74 (Only 21 CFR §§ 74.101-706), 81, 82 (Only 21 CFR §§ 82.3-82.706), 225, 226, 500.23, 500.24, 500.29, 500.45, 500.50, 500.80-500.92, 501, 502, 507, 509, 510.301, 510.305, 558, 570 (Except 21 CFR §§ 570.6, 570.15, and 570.17), 573, 579, 582, 584, and 589.
3. The Code of Federal Regulations, Title 21 CFR Part 1 Subpart L, Subpart M, and Subpart O.

**Authority: O.C.G.A. § 2-13-12.**