CLAIMS

1. Use of a peritoneal dialysate comprising 1.5 to 1.8 w/v% of glucose and 120 to 130 mEq/L of sodium in the manufacture of a medicament for peritoneal dialysis, wherein the peritoneal dialysate is for independent use once or twice a day, and

wherein the peritoneal dialysate is used in combination with another peritoneal dialysate that contains 1.35 to 2.5 w/v% of glucose and 132 to 135 mEq/L of sodium, in peritoneal dialysis that involves injection of an amount of 1.5 to 2.0 L per one time in an abdominal cavity of a subject, storage and drainage three to five times a day.

2. The use according to claim 1, wherein the peritoneal dialysate comprises 123 to 128 mEq/L of sodium.

3. The use according to claim 1 or 2, wherein the peritoneal dialysate further comprises 2.0 to 4.0 mEq/L of calcium; 0.5 to 1.5 mEq/L of magnesium; 86.0 to 98.0 mEq/L of chlorine; and 35 to 40 mEq/L of an alkalizing agent.

4. The use according to any one of claims 1 to 3, wherein the another peritoneal dialysate further contains 2.3 to 4.0 mEq/L of calcium; 0.5 to 1.5 mEq/L of magnesium; 95.0 to 105.5 mEq/L of chlorine; and 35 to 40 mEq/L of an alkalizing agent.

5. The use according to any one of claims 1 to 4, wherein the peritoneal dialysate and the another peritoneal dialysate are provided as a peritoneal dialysate set.