

PHARMATRANS SANAQ AG

PHARMACEUTICALS

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SPECIFICATION

SODIUM STARCH GLYOLATE SSG SANAQ

Expmt. Methods: USP / EP

TEST ITEM SPECIFICATIONS

Appearance White, relatively free-flowing powder

Identification

A: FT-IR Corresponds to the standard B: It is colored blue by Iodine

C: A white crystalline precipitate is formed

D: An intense yellow color to a nonluminous flame

pH Between 5.50 and 7.50
Loss on drying Not more than 10.0 %
Iron Not more than 0.002 %
Sodium Chloride Not more than 7.0 %
Assay (Sodium) 2.8 % ~ 4.2 % of Sodium
Residual Solvent (Methanol) Not more than 0.3 %

Residual Solvent (Methanol)
Heavy Metals
Sodium Glycolate

Not more than 0.3 %
Not more than 0.002 %
Not more than 2.0 %

Microbial Limits

Salmonella Species Absent Escherichia Coli Absent

Particle size

 \leq 40µm not less than 40% \geq 80µm not more than 7%