Succinylcholine Chloride

[ISUCC1] Relaxin®500mg/20mL/VialATC Code : M03AB01[I1SUCC] Relaxin®500mg/20mL/VialATC Code : M03AB01

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適應症: 手術用於鬆弛肌肉。

藥理分類: Neuromuscular Blocker Agent, Depolarizing.

用法用量: Administration: IV, IM.

Dosage regimens:

Adults:

-IV

Short procedures:

0.6 mg/kg IV (range 0.3-1.1 mg/kg) over 10-30 seconds.

Test Dose 0.1 mg/kg (5-10 mg). If test dose produces moderate muscle relaxation, 20 mg dose probably sufficient for short procedures; if test dose produces minimum relaxation, 30 mg probably needed.

Prolonged procedures:

0.3-1.1 mg/kg intermittent IV injection initially followed by 0.04-0.07 mg/kg at appropriate intervals. Alternatively, 0.5-10 mg/min by continuous IV infusion depending on the response.

-IM: up to 3-4 mg/kg, if suitable vein is inaccessible; MAX 150 mg.

Children:

—IV: infants & small children, 2 mg/kg; older children & adolescents, 1 mg/kg. Additional doses may be given if necessary.

Continuous IV infusions considered unsafe in neonates and children.

—IM: infants & older children, up to 3-4 mg/kg, if suitable vein is inaccessible; MAX 150 mg.

不良反應: 心跳停止、徐脈、頻脈、不整脈、低血糖、支氣管痙攣、遷延性無呼吸、惡性 過高熱、眼內壓上升、肌痛、發疹、過敏等。

注意事項: 1.Anticholinesterase 製劑會阻礙本劑分解。

2.若與 pH > 8.5 之鹼性溶液混合 (如 barbiturate solutions) 因會產生沈澱,故不得使用於同一注射筒內。

懷 孕 期: 1. 本藥應只使用於確切需要之病況。

2.Small amounts cross the placenta. Sensitivity to succinylcholine may be increased due to a ~24% decrease in plasma cholinesterase activity during pregnancy and several days postpartum.

授 乳 期: It is not known if succinylcholine is present in breast milk. The manufacturer recommends that caution be exercised when administering succinylcholine to breastfeeding women.

安 定 性: 配製後溶液冷藏(2-8℃)可保存24小時。

相容輸注液: NS、D5W、D10W、D5NS、1/2NS、Dextrose 5% in Lactated Ringer's、Lactated Ringer's、Ringer's inj.、Fructose 10% in NaCl 0.9%、Fructose 10% in water。

備 註: 批價代碼【IISUCC】為手術麻醉專用碼。