

## KREBS-RINGER SOLUTION FOR 5.5 L OF WATER (172 mM)

Prepare in this sequence:

1. Sodium Chloride 124	79.72g	
2. Potassium Chloride 2.5	2.05g	↙
3. Sodium Phosphate, monobasic 1.25	1.90g	↙
4. Sodium Bicarbonate 26	24.03g	USE FINE SCALES

Prepare separately 100ml of Stock 1M Calcium Chloride & 100 ml of Stock Magnesium Sulfate solutions in water:

1. Calcium Chloride	14.7g
2. Magnesium sulfate (w/7H <sub>2</sub> O)	24.65g

*Before preparing the solution:*

- Clean the scales surfaces with wet and then dry kimwipes
- Rinse weighing spatulas in deionized water and dry out with kimwipes
- Take weighing dishes only from closed boxes in the draw
- Put weighing dishes and weighing spatulas only on a clean paper towel
- After each chemical weighed wipe out the spatulas with kimwipes
- After weighing chemicals tightly close the cups of the chemicals/ use only separate bottles of these chemicals/ not on stock shelf
- After the solution is stirred well tightly close the cup on the bottle and put the bottle in the cold room right away
- Btl of cold water to make this up is in cold room

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For 1.5L of the final solution:

- 750ml Ringer +750ml deionized water
- + 3.0 ml of Stock Calcium Chloride (2mM)
- + 1.5 ml of Stock Magnesium Sulfate (1mM)
- + 1.5 scoop of Dextrose (10mM)

## Sucrose Solution

	250ml	500ml
5x Stock solution (ml)	50	100
DDW (ml)	200	400
Ca <sup>2+</sup> (0.5mM, ml)	0.12	0.24
Mg <sup>2+</sup> (7mM, ml)	1.7	3.4
Sucrose (gram)	19	38

From Geiger JR and Jonas P (2000) Dynamic control of presynaptic Ca<sup>2+</sup> inflow by fast-interactivating K<sup>+</sup> channels in hippocampal mossy fiber boutons. *Neuron* 28:927-939

For dissection and storage of slices, a sucrose-containing physiological saline with (**make 4 liters**)

NaCl	87 mM	20.34 g
Sucrose	75 mM	102.69 g
NaHCO <sub>3</sub>	25 mM	8.4 g
Glucose	25 mM	18.02 g
KCl	2.5 mM	0.745 g
<b>NaH<sub>2</sub>PO<sub>4</sub></b>	<b>1.25 mM</b>	<b>0.69</b>
CaCl <sub>2</sub>	0.5 mM	-from stock
MgCl <sub>2</sub>	7 mM	-from stock

Storage in 35<sup>0</sup>C, 1st 30min, use **sucrose** solution then transfer to normal solution for another 30min.