

[Protocol 1.02.1] Preparation of 0.5 mM EDTA

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<p>Protocol 1.02.1 Preparation of 0.5mM EDTA</p> <p>Version: 1.0 (20.12.2021)</p> <p>Media and Reagents:</p> <p>UltraPure™ 0.5 M EDTA (Thermo Fisher,15575-020)</p> <p>PBS w/o Ca2+, Mg2+ (Life Technologies, 14190250)</p> <p>Materials and Equipment:</p> <p>Pipettes</p> <p>Filtered pipette tips</p> <p>5 mL, 10 mL pipettes</p> <p>1. Introduction and Purpose</p> <p>This protocol describes how to aliquot/prepare EDTA according to the concentration, which is used for the chemical passage of induced pluripotent stem cells (iPSC) cultured in Geltrex or Matrigel coated well plates.</p> <p>2. Aliquoting 0.5 M EDTA</p> <p>Pipette 10 - 13 ml of UltraPure™ 0,5 M EDTA (pH 8.0) into a 15 ml Falcon tube and store this at room temperature on the rack in the cell culture lab</p> <p>3. Preparation of 0.5 mM EDTA</p> <p>1. Prepare 0.5 mM EDTA by adding the appropriate volume of UltraPure™ 0,5 M EDTA (pH 8.0) to PBS w/o Mg2+, Ca2+ (1:1000 dilution).</p> <p>2. Label it with the detailed name of the solution, e.g. 0.5 mM EDTA + PBS (-/-), the name of who prepared, and the date of the preparation.</p> <p>3. Store the labeled 15 ml Falcon tube with 0.5 mM EDTA at room temperature on the rack in the cell culture lab</p>
