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## 🌐 Electrophysiology with iPSC-derived neurons

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### ABSTRACT

This protocol describes the method to perform patch-clamp recordings of iPSC-derived neurons.

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**Protocol status:** Working

We use this protocol and it's working. This protocol was originally provided by Olga Kopach.

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- 1 Patch-clamp recordings of iPSC-derived neurons are performed using an infrared differential interference contrast (DIC) imaging system on an Olympus BX51WI upright microscope (Olympus, Japan) coupled with a Multipatch 700B amplifier under the control of pClamp 10.2 software package (Molecular Devices, USA)
- 2 For the recordings, a neuronal culture or co-culture is plated on glass coverslips, placed in a recording chamber mounted on the microscope stage and constantly perfused with a physiological buffer medium.
- 3 Whole-cell recordings are performed using glass pipettes with a resistance of 3.5-6 MΩ when filled with the intracellular solution.

#### Note

##### Perfusion medium (in mM):

126 NaCl, 3 KCl, 2 MgSO<sub>4</sub>, 2 CaCl<sub>2</sub>, 26 NaHCO<sub>3</sub>, 1.25 NaH<sub>2</sub>PO<sub>4</sub>, 10 D-glucose

The medium is continuously bubbled with. [M] 95 % volume O<sub>2</sub> and [M] 5 % volume CO<sub>2</sub> (pH 7.4) and maintained at 30 °C - 33 °C.

- 4 Whole-cell recordings are performed using glass pipettes with a resistance of 3.5-6 MΩ when filled with the intracellular solution.

#### Note

##### Intracellular solution (in mM):

126 K-gluconate, 4 KCl, 4 MgCl<sub>2</sub>, 2 BAPTA, 4 Mg-ATP, 0.4 Na-ATP (pH 7.2, osmolarity ~295 mOsmol)

- 5 In the whole-cell (immediately after membrane breakthrough), iPSC-derived neurons are recorded for the resting membrane potential ( $V_{rest}$ ), membrane capacitance ( $C_m$ ), the membrane time constant ( $\tau_m$ ), and input resistance ( $R_{in}$ )
- 6 To induce neuronal firing, a series of sub- and supra-threshold rectangular current pulses are applied with a stepwise-increased stimulus intensity at the  $V_{hold}$  set at  $-60$  mV to  $-70$  mV.

#### Note

The second protocol tested is a slow-ramp current injection, ramped up with a  $100$ – $200$  pA/s slope.

- 7 The analysis of the AP waveform was performed for the first AP only to quantify the threshold value, the spike amplitude, overshoot, the spike width (duration at half-maximal amplitude), the rates of depolarisation and re-polarisation phase.