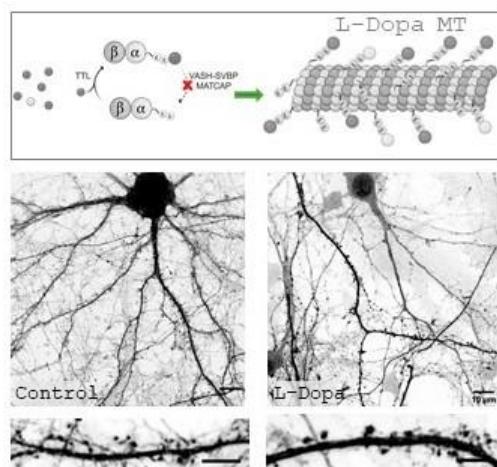


L-DOPA INCORPORATION INTO TUBULIN ALTERS MICROTUBULE DYNAMICS AND REDUCES DENDRITIC SPINE INVASION AND SYNAPSE MAINTENANCE

Previous research demonstrated that L-dopa, a tyrosine analog used in the treatment of Parkinson's disease, can be incorporated into α -tubulin and form microtubules. Now we show that L-dopa treatment in mature wild-type hippocampal neurons reduces dendritic spine density and the number of excitatory synapses, affecting the quality and quantity of synapses. The incorporation of L-dopa into α -tubulin disrupts microtubule dynamics, leading to decreased microtubule invasion into dendritic spines and increased synaptotoxicity. In neurons lacking key enzymes in the tubulin tyrosination/detyrosination cycle, these effects are absent, confirming the role of L-dopa-modified tubulin in synaptic dysfunction. This highlights the importance of balanced tubulin modification in synaptic health and the need for new strategies in Parkinson's treatment to avoid such side effects.



JUEVES 5 de SEPTIEMBRE . 12:00 h

Zoom: <https://zoom.us/j/91097843541?pwd=alVVeXBQZDlrOE5NYW53dVh0WXc3UT09>

ID de reunión: 910 9784 3541 **Código de acceso:** 489930



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INVITAN DENTRO DE SUS CICLOS DE SEMINARIOS 2024