



MISSOURI HOUSE OF REPRESENTATIVES
WITNESS APPEARANCE FORM

BILL NUMBER: HB 2922		DATE: 3/9/2026	
COMMITTEE: Emerging Issues			
TESTIFYING: <input checked="" type="checkbox"/> IN SUPPORT OF <input type="checkbox"/> IN OPPOSITION TO <input type="checkbox"/> FOR INFORMATIONAL PURPOSES			
WITNESS NAME			
INDIVIDUAL:			
WITNESS NAME: AIDAN EVERETT WAGNER		PHONE NUMBER:	
BUSINESS/ORGANIZATION NAME:		TITLE:	
ADDRESS:			
CITY:		STATE:	ZIP:
EMAIL:	ATTENDANCE: Written		SUBMIT DATE: 3/9/2026 8:09 AM
THE INFORMATION ON THIS FORM IS PUBLIC RECORD UNDER CHAPTER 610, RSMo.			



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WITNESS NAME			
INDIVIDUAL:			
WITNESS NAME: ARNIE C. "HONEST-ABE" DIENOFF-STATE PUBLIC ADVOCAT		PHONE NUMBER:	
BUSINESS/ORGANIZATION NAME:		TITLE:	
ADDRESS:			
CITY:		STATE:	ZIP:
EMAIL:	ATTENDANCE: In-Person		SUBMIT DATE: 3/9/2026 11:55 PM
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I am in favor of Access to Alternate Therapies.



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WITNESS NAME		
INDIVIDUAL:		
WITNESS NAME: SUZANNE OPPERMAN, PMHNP CCTP-II CDAC		PHONE NUMBER:
BUSINESS/ORGANIZATION NAME:		TITLE:
ADDRESS:		
CITY:		STATE: ZIP:
EMAIL:	ATTENDANCE: Written	SUBMIT DATE: 3/6/2026 1:38 PM

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Severe Chronic PTSD is a severely debilitating illness. This bill would allow providers to have assess to treat treatment resistant clients with drugs such as Ibodine, MDMA (Midomafetarmine), TSND-201 (Methylone), Psilocybin, Ketamine and others. As a Veteran and Mental Health Trauma and addiction specialist, Clients would hugely benefit from many medications not available at this time.

These severely affected clients become totally dependent on Medicaid treatment, illegal substances, unable to work, and often end up incarcerated due to their severe mental illness



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WITNESS NAME			
INDIVIDUAL:			
WITNESS NAME: SARAH BERRY		PHONE NUMBER:	
BUSINESS/ORGANIZATION NAME:		TITLE:	
ADDRESS:			
CITY:		STATE:	ZIP:
EMAIL:	ATTENDANCE: Written	SUBMIT DATE: 3/9/2026 9:34 AM	
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Missouri already operates under the federal Right-to-Try Act.

So the real question for this committee is simple:

What problem does HB 2922 solve that federal law does not already address?

This bill primarily expands liability protections for manufacturers and prescribers while allowing access to drugs that have only completed Phase I safety testing — treatments whose effectiveness is still unknown.

When a policy both weakens accountability and shifts financial risk to patients, lawmakers should ask whether the bill is truly expanding patient choice or simply expanding legal protection for the companies involved.