Pronovost-Morgan C, Greenway KT, Roseman L, The ReSPCT Experts. An international Delphi consensus for reporting of setting in psychedelic clinical trials. *Nat Med.* 2025. https://doi.org/10.1038/s41591-025-03685-9

Item	Item Number	Item details	Reported on page number:
Physical environment			
Location	1	Location of the trial, and whether indoors/outdoors and urban/rural/suburban.	
sAmbiance	2	Room ambiance curated by the study team.	
Access to nature	3	Sources of nature or natural elements that are physically or visually accessible to participants.	
Objects and decorations	4	Objects and decorations in the room.	
Lighting	5	Room lighting and adjustability.	
Sensory reduction	6	Sensory reduction used, such as headphones and eyeshades.	
Bathroom facilities	7	Level of bathroom accessibility and privacy.	
Dosing session procedure			
Number and roles of people present	8	Number and roles of people present, including participants, study staff and informal support.	
Positioning	9	Relative position of people in the room, and what participants were positioned on (e.g., bed, mat, couch).	
Focus and main activities	10	Focus (internal or external) and main activities of the dosing session.	
Music or soundscapes	11	Music or soundscapes that accompanied the dosing experience.	
Interpersonal interventions	12	Verbal or physical interpersonal interventions used throughout the session, and how consent was obtained.	
Participant autonomy, control, and agency	13	Level of participant control and agency over activities and environment of the dosing session.	
Dosing regimen	14	Dosing regimen, including drug dose(s), frequency, route of administration and length of the dosing session.	
Medical and experimental procedures or assessments	15	Medical and experimental procedures or assessments performed during the dosing session.	
Pre- and post-dosing protocol	16	Activities that took place immediately prior to- or post-dosing, including participant arrival and release conditions.	
Potential disturbances or interruptions	17	Disturbances or interruptions that may have impacted the quality of the dosing session.	
Therapeutic framework and protocol			
Therapeutic or guiding approach	18	Therapeutic/guiding approach used throughout the study, if any, with the accompanying manual or protocol.	
Narrative framing	19	Framing of the trial intervention by the study team, including the short- and long-term drug effects.	
Number of sessions	20	Number and length of preparation, dosing, and integration sessions.	

Preparation protocol	21	Activities performed during the preparation sessions.	
Integration protocol	22	Activities performed during the integration sessions.	
Additional support/follow-up	23	Formal or informal support or follow-up offered to participants after the end of the trial.	
Study personnel qualifications	24	The credentials, training, and expertise of personnel providing the study intervention or care.	
Cultural competence and safety	25	Study team's level of cultural competence and efforts towards cultural safety.	
Subjective experiences			
Therapeutic alliance	26	Therapeutic alliance between participants and facilitators throughout the intervention.	
Trust	27	Participant's level of trust throughout the intervention.	
Physical comfort	28	Participant's level of physical comfort during the dosing session.	
Physical safety	29	Participant's sense of physical safety during the dosing session.	
Psychological and cultural safety	30	Participant's sense of interpersonal safety with the people present during the session.	