Rev. 06/20

IDPH New Medical Cannabidiol Product Form
Laboratory Verification and Validation Studies

In the development of any new product, prior to sampling and delivering a new product to a laboratory for a verification or validation study, the following form must be completed by a manufacturer, laboratory, and IDPH.

Page 1 to be completed by a manufacturer

Manufacturer	Process Lot (concentrate)	Description of process lot:		
		Solvent(s) used:		
		Additives (carrier oils, terpenes, etc.):		
	Package Lot (Final Product)	Description of Product Form:		
		Product Matrix Ingredients:		
		THC/THCa concentration (range)	Per Dose	Per Unit
		CBD/CBDa concentration (range)	Per Dose	Per Unit
	Reference Material/ Documentation	Please describe the reference material and/or documentation the manufacturer intends to provide a laboratory to ensure a thorough and accurate verification or validation study.		
	Manufacturer Comments:			

Manufacturer Authorized Signature: _____

Date: _____

IDPH
IOWA Department of PUBLIC HEALTH

Rev. 06/20

Page 2 to be completed by a Laboratory

LAB	Identify validation / verification requirement	Validation or Verification
	Identify specific list of items needed from manufacturer:	
	Identify timeline for completion:	
	Laboratory Comments:	

Laboratory Authorized Signature: _____

Date: _____

Rev. 06/20

Page 3 to be completed by IDPH

Please refer to v4.2 Laboratory Acceptance Criteria Document for information pertaining to type and number of tests.

IDPH	Identify analytes required for testing, as well as standard or reduced testing.	Potency	Total aerobic microbial count
		Metals	Total combined yeast and mold count
		Pesticides	Aspergillus
		Solvents:	Shiga-Toxin Producing E.coli
		Other:	*Salmonella*
	IDPH Comments:		

IDPH Authorized Signature: _____

Date: _____