

State Medical Board of Ohio Report of RU-486 Event

(Required pursuant to R.C. 2919.123)

To be completed by the physician who provided RU-486

1. Date RU-486 was provided:	<u> MAY </u>	<u> 07 </u>	<u> 2018 </u>
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided:	Women's Med Dayton		
3. Address of medical practice or facility at which RU-486 was provided:	1401 E Stroop Rd Dayton, Ohio 45429		
4. Date post RU-486 complication began:	<u> 05/15/18 </u>		
5. Event(s) (Please check all that apply):	<input checked="" type="checkbox"/> Incomplete abortion <input type="checkbox"/> Adverse reaction to RU-486 <input type="checkbox"/> Patient hospitalized <input type="checkbox"/> Patient received a transfusion <input type="checkbox"/> Severe bleeding <input type="checkbox"/> Other: serious event (specify) _____		
6. Duration of event:	<u> 7 </u> Hours	<u> </u> Days	
7. Remarks:			
8. a. Name of physician who provided RU-486	<u> D. K. K... MD </u>		
8. b. Physician's signature	<u> D. K. K... </u>	<u> MD, D.O. </u>	
	Date	<u> 5/15/18 </u>	

Send completed forms to: State Medical Board of Ohio
Legal Department
30 E. Broad St., 3rd Floor
Columbus, OH 43215-6127

MEDICAL BOARD
MAY 21 2018