



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:	11	04	2015
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: <i>Planned Parenthood SW Ohio Region</i>			
3. Address of medical practice or facility at which RU-486 was provided: <i>2314 Auburn Ave, Cincinnati, OH 45219</i>			
4. Date post RU-486 complication began: <i>11/20/15</i>			
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: <i>1</i> Hours _____ Days			
7. Remarks: <i>D+C performed w/o incident.</i>			
8. a. Name of physician who provided RU-486 _____ <i>Dr. [Signature]</i>			
8. b. Physician's signature _____ <i>[Signature]</i> <i>MD/DO</i>			
Date <i>12/4/15</i>			

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

MEDICAL BOARD
DEC 09 2015



State Medical Board of Ohio Report of RU-486 Event

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To be completed by the physician who provided RU-486

1. Date RU-486 was provided:	<u>6</u>	<u>2</u>	<u>15</u>	
	Month	Day	Year	
2. Name of medical practice or facility at which RU-486 was provided: <u>Planned Parenthood Southwest Ohio</u>				
3. Address of medical practice or facility at which RU-486 was provided: <u>2314 Auburn Ave Cincinnati, OH 45219</u>				
4. Date post RU-486 complication began: <u>6/18/15</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: _____ Hours <u>30</u> Days <u>Follow up period after mib</u>				
7. Remarks: <u>pt. started to attempt completion with second dose of miso prostol, had D+C on 7/21/15 without problem</u>				
8. a. Name of physician who provided RU-486 <u>Charon King</u>				
8. b. Physician's signature _____ M.D./D.O. _____				
Date <u>7/24/15</u>				

MEDICAL BOARD
AUG 3 2015

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State Medical Board of Ohio Report of RU-486 Event

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To be completed by the physician who provided RU-486

1. Date RU-486 was provided:	<u>2</u>	<u>17</u>	<u>2015</u>
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided:	<u>Planned Parenthood Southwest Ohio Region</u>		
3. Address of medical practice or facility at which RU-486 was provided:	<u>2314 Auburn Ave.</u>		
4. Date post RU-486 complication began:	<u>3/6/15</u>		
5. Event(s) (Please check all that apply):	MEDICAL BOARD		
<input checked="" type="checkbox"/> Incomplete abortion	<input type="checkbox"/> Adverse reaction to RU-486	<input type="checkbox"/> Patient hospitalized	AUG 3 2015
<input type="checkbox"/> Patient received a transfusion	<input type="checkbox"/> Severe bleeding		
<input type="checkbox"/> Other serious event (specify) _____			
6. Duration of event: _____ Hours <u>14</u> Days <u>Follow up period after medo.</u>			
7. Remarks:	<u>Pt. did well with second dose of misoprostol</u>		
8. a. Name of physician who provided RU-486	<u>Dr. Kelsey</u>		
8. b. Physician's signature	<u>[Signature]</u>	<u>M.D./D.O.</u>	
	Date _____		

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