

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>10</u>	<u>27</u>	<u>2017</u>
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: <u>Preterm</u>			
3. Address of medical practice or facility at which RU-486 was provided: <u>12000 Shaker Blvd Cleveland, Ohio 44120</u>			
4. Date post RU-486 complication began: <u>11/29/17</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>3</u> Hours _____ Days			
7. Remarks: 			
8. a. Name of physician who provided RU-486 <u>Mitchell Reider, MD</u>			
8. b. Physician's signature <u>[Signature]</u> MD/DO _____			
Date <u>12/1/17</u>			

Send completed forms to:

State Medical Board of Ohio

Legal Department

30 E. Broad St., 3rd Floor

Columbus, OH 43215-6127


MEDICAL BOARD

DEC 05 2017

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:	10	28	2017
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: Preterm			
3. Address of medical practice or facility at which RU-486 was provided: 12000 Shaker Blvd. Cleveland, OH 44120			
4. Date post RU-486 complication began: 11/11/17			
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: 3 Hours _____ Days			
7. Remarks:			
8. a. Name of physician who provided RU-486 Justin Lappen, MD			
8. b. Physician's signature  MD/D.O.			
Date 12/2/17			

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MEDICAL BOARD

DEC 07 2017

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>10</u>	<u>28</u>	<u>2017</u>
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: <u>Preterm</u>			
3. Address of medical practice or facility at which RU-486 was provided: <u>12000 Shaker Blvd. Cleveland, OH 44120</u>			
4. Date post RU-486 complication began: <u>11/17/2017</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>2</u> Hours _____ Days			
7. Remarks: 			
8. a. Name of physician who provided RU-486 <u>Justin Lappen, MD</u>			
8. b. Physician's signature <u>[Signature]</u> M.D./D.O. Date <u>12/2/17</u>			

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MEDICAL BOARD

DEC 07 2017



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: 7 / 12 / 17
Month Day Year

2. Name of medical practice or facility at which RU-486 was provided:
Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:
25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began: 7/19/17

5. Event(s) (Please check all that apply):
☒ Incomplete abortion ☐ Adverse reaction to RU-486 ☐ Patient hospitalized
☐ Patient received a transfusion ☐ Severe bleeding
☐ Other serious event (specify) _____

6. Duration of event: _____ Hours _____ Days

7. Remarks: Med abortion process was initiated per FDA regimen on 7/12/17. Followup ultrasound on 7/19/17. Showed an ongoing pregnancy. Surgical aspiration was done on 7/20/17, pt did well post-op.

8. a. Name of physician who provided RU-486 Timothy Kress, MD

8. b. Physician's signature Timothy Kress MD/DO
Date 8/15/17

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30 E. Broad St., 3rd Floor
Columbus, OH 43215-6127



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>7</u> / <u>13</u> / <u>17</u> Month / Day / Year
2. Name of medical practice or facility at which RU-486 was provided:	<u>Planned Parenthood of Greater Ohio</u>
3. Address of medical practice or facility at which RU-486 was provided:	<u>25350 Rockside Rd. Bedford Heights, Ohio 44146</u>
4. Date post RU-486 complication began:	<u>7/17/17</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>1</u> Hours <u> </u> Days
7. Remarks:	<u>Med abortion process initiated per FDA regimen on 7/13/17. Second dose of misoprostol was given on 7/17/17 due to lack of results from first dose. Follow-up ultrasound on 7/19/17 showed ongoing pregnancy. Surgical aspiration was performed and pt did well post-op.</u>
8. a. Name of physician who provided RU-486	<u>TIMOTHY KRESS, MD</u>
8. b. Physician's signature	<u>Timothy Kress</u> <u>MD/DO</u>
Date	<u>8/15/17</u>

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Columbus, OH 43215-6127

MEDICAL BOARD

AUG 23 2017



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:

7 / 19 / 17
Month Day Year

2. Name of medical practice or facility at which RU-486 was provided:

Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:

25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began:

8/2/17

5. Event(s): (Please check all that apply):

☒ Incomplete abortion ☐ Adverse reaction to RU-486 ☐ Patient hospitalized

☐ Patient received a transfusion ☐ Severe bleeding

☐ Other serious event (specify) _____

6. Duration of event: 1 Hours _____ Days

7. Remarks: Medication abortion procedure was initiated per FDA regimen on 7/19/17. At follow-up visit on 8/2/17 ultrasound revealed continued pregnancy. Surgical abortion was performed the same day and pt. did well post-op.

8. a. Name of physician who provided RU-486

Timothy Kress, MD

8. b. Physician's signature

Timothy Kress MD/DO

Date

9/1/17

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Columbus, OH 43215-6127

MEDICAL BOARD

SEP 15 2017



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:

Month 8 Day 1 Year 17

2. Name of medical practice or facility at which RU-486 was provided:

Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:

25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began:

8/15/17

5. Event(s) (Please check all that apply):

☒ Incomplete abortion

☐ Adverse reaction to RU-486

☐ Patient hospitalized

☐ Patient received a transfusion

☐ Severe bleeding

☐ Other serious event (specify) _____

6. Duration of event: 1 Hours Days

7. Remarks: Medication abortion procedure was initiated per FDA regimen on 8/1/17. At follow-up visit on 8/15/17, ultrasound revealed a continuing pregnancy. Surgical abortion was done the same day and pt. did well post-op.

8. a. Name of physician who provided RU-486

Timothy Kress, MD

8. b. Physician's signature

Timothy Kress

MD/DO

Date

9/1/17

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MEDICAL BOARD

SEP 15 2017



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:	8	1	17
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: Planned Parenthood of Greater Ohio			
3. Address of medical practice or facility at which RU-486 was provided: 25350 Rockside Rd. Bedford Heights, Ohio 44146			
4. Date post RU-486 complication began: 8/18/17			
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 _____ Days			
7. Remarks: Med abortion procedure was initiated per FDA regimen on 8/1/17. Bloodwork on 8/16/17 and ultrasound on 8/22/17 revealed an incomplete abortion. Pt chose to repeat the medication regimen on 8/22/17; ultrasound on 8/29/17 showed abortion was complete.			
8. a. Name of physician who provided RU-486: Timothy Kress MD			
8. b. Physician's signature: <u>Timothy Kress</u> (MD) / D.O. Date: 9/1/17			

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SEP 15 2017



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To be completed by the physician who provided RU-486

1. Date RU-486 was provided:

8 / 2 / 17
Month Day Year

2. Name of medical practice or facility at which RU-486 was provided:

Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:

25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began: 10/3/17

5. Event(s) (Please check all that apply):

☒ Incomplete abortion

☐ Adverse reaction to RU-486

☐ Patient hospitalized

☐ Patient received a transfusion ☐ Severe bleeding

☐ Other serious event (specify) _____

6. Duration of event: 1 Hours Days

7. Remarks: Med ab procedure started 8/2/17 per FDA protocol. Pt. returned for follow-up on 10/3/17 at which time ultrasound showed a continuing pregnancy. Surgical abortion was performed on 10/14/17; pt did well post-op.

8. a. Name of physician who provided RU-486 TIMOTHY KRESS, MD

8. b. Physician's signature

Timothy Kress MD/DO

Date 11/10/2017

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MEDICAL BOARD

NOV 17 2017



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: 8 / 9 / 17
Month Day Year

2. Name of medical practice or facility at which RU-486 was provided:
Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:
25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began: 8/15/17

5. Event(s) (Please check all that apply):

☒ Incomplete abortion

☐ Adverse reaction to RU-486

☐ Patient hospitalized

☐ Patient received a transfusion ☐ Severe bleeding

☐ Other serious event (specify) _____

6. Duration of event: 1 Hours _____ Days

7. Remarks: Medication abortion procedure was initiated per FDA regimen on 8/9/17. At follow-up visit on 8/15/17, ultrasound revealed a continuing pregnancy. Surgical abortion was done on 8/17/17 and the pt. did well post-op.

8. a. Name of physician who provided RU-486 _____

8. b. Physician's signature _____

Date _____

MEDICAL BOARD
7/5/2017
OCT 04 2017

Andrey G. Fiksel, MD

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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:

8 / 15 / 17
Month Day Year

2. Name of medical practice or facility at which RU-486 was provided:

Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:

25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began:

8/24/17

5. Event(s) (Please check all that apply):

☒ Incomplete abortion

☐ Adverse reaction to RU-486

☐ Patient hospitalized

☐ Patient received a transfusion ☐ Severe bleeding

☐ Other serious event (specify) _____

6. Duration of event: 1 Hours _____ Days

7. Remarks: Medication abortion procedure was initiated per FDA regimen on 8/15/17. At follow-up visit on 8/24/17 ultrasound showed absence of gestational sac, but incomplete abortion. Surgical aspiration was done on 8/24/17 and pt. did well post op.

8. a. Name of physician who provided RU-486

Timothy Kress, MD

8. b. Physician's signature

Timothy Kress, MD

MD/DO

Date

9/11/17

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MEDICAL BOARD

SEP 15 2017



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: 9 / 8 / 17
Month Day Year

2. Name of medical practice or facility at which RU-486 was provided:
Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:
25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began: 10/14/17

5. Event(s) (Please check all that apply):

☒ Incomplete abortion ☐ Adverse reaction to RU-486 ☐ Patient hospitalized

☐ Patient received a transfusion ☐ Severe bleeding

☐ Other serious event (specify) _____

6. Duration of event: 1 Hours Days

7. Remarks: med ab procedure started 9/8/17 per FDA protocol. Pt. returned for follow-up on 10/14/17 at which time ultrasound showed a continuing pregnancy. Surgical abortion was performed on 10/14/17; pt did well post-op.

8. a. Name of physician who provided RU-486 TIMOTHY KRESS, M.D.

8. b. Physician's signature Timothy Kress M.D./D.O.

Date 11/10/2017

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MEDICAL BOARD

NOV 17 2017



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:	9	8	17
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: Planned Parenthood of Greater Ohio			
3. Address of medical practice or facility at which RU-486 was provided: 25350 Rockside Rd. Bedford Heights, Ohio 44146			
4. Date post RU-486 complication began: 9/19/17			
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: 1 Hours _____ Days			
7. Remarks: Medication ab procedure started on 9/8/17. Follow-up bloodwork + ultrasound showed continuing pregnancy. Surgical aspiration completed on 9/23/17; pt did well post-op.			
8. a. Name of physician who provided RU-486: Timothy Kress, MD			
8. b. Physician's signature: <u>Timothy S. Kress</u> MD/DO			
Date: 10/6/17			

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OCT 20 2017

MEDICAL BOARD



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:	9	13	17
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: Planned Parenthood of Greater Ohio			
3. Address of medical practice or facility at which RU-486 was provided: 25350 Rockside Rd. Bedford Heights, OH 44146			
4. Date post RU-486 complication began: 10/3/17			
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: 1 Hours _____ Days			
7. Remarks: Med abortion procedure started 9/13/17 per FDA protocol. Follow-up bloodwork on 10/3/17 indicated an incomplete abortion and ultrasound showed a continuing pregnancy. Surgical abortion was performed on 10/31/17, pt did well post-op			
8. a. Name of physician who provided RU-486: TIMOTHY KRESS, MD.			
8. b. Physician's signature: <u>Timothy Kress</u> (MD) / D.O. Date: 11/10/2017			

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MEDICAL BOARD

NOV 17 2017



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:	9	27	17
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: Planned Parenthood of Greater Ohio			
3. Address of medical practice or facility at which RU-486 was provided: 25350 Rockside Rd. Bedford Heights, Ohio 44146			
4. Date post RU-486 complication began: 10/11/17			
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 _____ Days			
7. Remarks: Med ab procedure started on 9/27/17 per FDA protocol. Follow-up ultrasound on 10/11/17 showed intrauterine debris. Surgical aspiration was done on 10/11/17; pt did well post-op.			
8. a. Name of physician who provided RU-486: TIMOTHY KRESS, MD			
8. b. Physician's signature: <u>Timothy Kress</u> MD/DO			
Date: 11/10/2017			

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MEDICAL BOARD

NOV 17 2017



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:

10
Month

21
Day

17
Year

2. Name of medical practice or facility at which RU-486 was provided:

Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:

25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began:

11/10/17

5. Event(s) (Please check all that apply):

☒ Incomplete abortion

☐ Adverse reaction to RU-486

☐ Patient hospitalized

☐ Patient received a transfusion ☐ Severe bleeding

☐ Other serious event (specify) _____

6. Duration of event: _____ Hours _____ Days

7. Remarks: Med ab process was initiated on 10/21/17, following FDA criteria. Follow up ultrasound on 11/10 showed continuing pregnancy. Surgical ab was done on 11/16/17 and pt. did well post-op.

8. a. Name of physician who provided RU-486

8. b. Physician's signature

Date

12/5/17

Send completed forms to:

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Columbus, OH 43215-6127

MEDICAL BOARD

DEC 21 2017



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: 12 / 9 / 17
Month Day Year

2. Name of medical practice or facility at which RU-486 was provided:
Planned Parenthood of Greater Ohio

3. Address of medical practice or facility at which RU-486 was provided:
25350 Rockside Rd. Bedford Heights, Ohio 44146

4. Date post RU-486 complication began: 12/29/17

5. Event(s) (Please check all that apply):
☒ Incomplete abortion ☐ Adverse reaction to RU-486 ☐ Patient hospitalized
☐ Patient received a transfusion ☐ Severe bleeding
☐ Other serious event (specify) _____

6. Duration of event: _____ Hours _____ Days

7. Remarks: Med ab procedure was initiated on 12/9/17 per FDA regimen. Pt returned for flu ultrasound on 12/29/17 and uterine debris was noted. Surgical aspiration was performed at that time; pt did well post op.

8. a. Name of physician who provided RU-486 Dr. T. Kress MD

8. b. Physician's signature T. Kress MD
Date 1/5/18

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Columbus, OH 43215-6127

MEDICAL BOARD

JAN 17 2018

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>7</u>	<u>10</u>	<u>2017</u>
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: <u>Northeast Ohio Women's Center</u>			
3. Address of medical practice or facility at which RU-486 was provided: <u>2127 State Rd Cuyahoga Falls Ohio 44223</u>			
4. Date post RU-486 complication began: <u>7/27/17</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 _____ Days			
7. Remarks: <u>post med AB patient had a remaining gestational sac w/ no fetal pole developed or remaining. patient had DEC on 7/27/17 to complete her process.</u>			
8. a. Name of physician who provided RU-486 <u>Dr. DAVID Burkons</u>			
8. b. Physician's signature <u>[Signature]</u> <u>MD/DO</u>			
Date <u>9/7/17</u>			

Send completed forms to: State Medical Board of Ohio

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MEDICAL BOARD

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>7</u>	<u>13</u>	<u>2017</u>
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: <u>Northeast Ohio Women's Center</u>			
3. Address of medical practice or facility at which RU-486 was provided: <u>2127 State Rd Cuyahoga Falls 44223</u>			
4. Date post RU-486 complication began: <u>8/12/17</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 _____ Days			
7. Remarks: <u>pt. had (+) pta post medication Abortion. She had heavy bleeding and on ultrasound there was remaining tissue but pregnancy was resolved. pt had a DEC on 8/22/17</u>			
8. a. Name of physician who provided RU-486: <u>Dr. L.A. Nunnally</u>			
8. b. Physician's signature: <u>[Signature]</u> MD/DO			
Date: <u>9/7/17</u>			

Send completed forms to: State Medical Board of Ohio

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Columbus, OH 43215-6127

MEDICAL BOARD

SEP 15 2017

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>October</u> <u>21</u> <u>2017</u> Month Day Year
2. Name of medical practice or facility at which RU-486 was provided:	<u>North east Ohio Women's Center</u>
3. Address of medical practice or facility at which RU-486 was provided:	<u>2127 State St</u> <u>Cuyahoga Falls, Ohio 44223</u>
4. Date post RU-486 complication began:	<u>11/16/17</u>
5. Event(s) (Please check all that apply):	<input checked="" type="checkbox"/> <u>Failed</u> 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>2</u> Hours <u> </u> Days
7. Remarks:	<u>a suction D+E was performed 11/16/17</u> <u>without difficulty</u>
8. a. Name of physician who provided RU-486	<u>Jennifer Watson</u>
8. b. Physician's signature	<u>J. Watson</u> MD/DO
Date	<u>11/18/17</u>

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MEDICAL BOARD

NOV 27 2017