

# Data Extraction Template

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## Item Settings

## Editor

[I want to start from scratch](#)

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H1

General information

+

Aa

Study ID

+

Aa

Title

+

Aa

Lead author contact details

+

🎯

Country in which the study conducted

1. –  
United States

2. –  
UK

3. –  
Canada

4. –  
Australia

5. –  
Other

+

Aa

Notes

H<sub>1</sub>

Characteristics of included studies

H<sub>2</sub>

Methods

Aa

Aim of study



Study design

1. –  
Randomised controlled trial
2. –  
Non-randomised experimental study
3. –  
Cohort study
4. –  
Cross sectional study
5. –  
Case control study
6. –  
Systematic review
7. –  
Qualitative research
8. –  
Prevalence study
9. –  
Case series
10. –  
Case report
11. –  
Diagnostic test accuracy study
12. –  
Clinical prediction rule
13. –

Economic evaluation

14. –

Text and opinion

15. –

Other



Aa

Start date



Aa

End date



Aa

Study funding sources



Aa

Possible conflicts of interest for study authors



H2

Participants



Aa

Population description



Aa

Inclusion criteria



Aa

Exclusion criteria



Target

Method of recruitment of participants

1. –

Phone

2. –

Mail

3. –

Clinic patients

4. –

Voluntary

5. –

Other



Total number of participants



Type of Mole

1. –

Complete Mole

2. –

Partial Mole

3. –

Invasive Mole

4. –

Choriocarcinoma

5. –

Placental Site Trophoblastic Tumor

6. –

Epithelioid Trophoblastic Tumor

7. –

Other



Time of Contraception Start

1. –

Immediately following GTD treatment

2. –

After beta-hCG normalization

3. –

Other



Baseline Characteristics



Contraception



Beta-hCG Normalization



Post-molar GTN



Incident Pregnancies



Adverse Events

## Preview

### General information

Study ID

Title

Title of paper / abstract / report that data are extracted from

Lead author contact details

Country in which the study conducted

- 1. ☐ United States
- 2. ☐ UK
- 3. ☐ Canada
- 4. ☐ Australia

5. ☐ Other

Notes

### Characteristics of included studies

## Methods

Aim of study

Study design

1. ☐ Randomised controlled trial
2. ☐ Non-randomised experimental study
3. ☐ Cohort study
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10. ☐ Case report
11. ☐ Diagnostic test accuracy study
12. ☐ Clinical prediction rule
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14. ☐ Text and opinion
15. ☐ Other

Start date

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Study funding sources

Possible conflicts of interest for study authors

## Participants

Population description

Inclusion criteria

Exclusion criteria

Method of recruitment of participants

1.

☐

Phone
2.

☐

Mail
3.

☐

Clinic patients
4.

☐

Voluntary
5.

☐

Other

Total number of participants

- Type of Mole
1.

☐

Complete Mole
2.

☐

Partial Mole
3.

☐

Invasive Mole
4.

☐

Choriocarcinoma
5.

☐

Placental Site Trophoblastic Tumor
6.

☐

Epithelioid Trophoblastic Tumor
7.

☐

Other

- Time of Contraception Start
1.

☐

Immediately following GTD treatment
2.

☐

After beta-hCG normalization
3.

☐

Other

Baseline Characteristics										
	Contraception Type 1			Contraception Type 2			Total Population		Additional Notes	
Contraception Type										
Age (mean)										
Gravidity (mean)										
Parity (mean)										
Contraception										
	Combined oral contraceptive pills	Vaginal Ring	Patch	Progestin only pills	Copper IUD	Progestin IUD	DMPA	Implant	Barrier Methods	Other
Number of Participants on Method										
Length of Time on Method (Average)										
Beta-hCG Normalization										

**Mean Time (in days) SD****Combined oral contraceptive pills****Vaginal Ring****Patch****Progestin only pills****Copper IUD****Progestin IUD****DMPA****Implant****Barrier Methods****Other****None**

Post-molar GTN

**Number of Participants****Combined oral contraceptive pills****Vaginal Ring****Patch****Progestin only pills****Copper IUD****Progestin IUD****DMPA****Implant****Barrier Methods****Other****None**

Incident Pregnancies

**Number of Participants****Combined oral contraceptive pills****Vaginal Ring****Patch****Progestin only pills****Copper IUD****Progestin IUD****DMPA****Implant**



Barrier Methods		Number of Participants					
Other							
None							
Adverse Events							
		Adverse Event Type 1	Number of Participants	Adverse Event Type 2	Number of Participants	Adverse Event Type 3	Number of Participants
Combined oral contraceptive pills							
Vaginal Ring							
Patch							
Progestin only pills							
Copper IUD							
Progestin IUD							
DMPA							
Implant							
Barrier Methods							
Other							
None							

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