

GRADE: Le Tappe del Processo

Definire il problema

Definire l'importanza relativa degli esiti

Ricerca le prove di effetto (RS, RCT)

Valutare la qualità delle prove per ciascun esito

Riassumere le prove per ciascun outcome

Valutare la qualità globale delle prove

Fare un bilancio dei benefici e degli eventi avversi

Definire la forza della raccomandazione

IMPLEMENTAZIONE E VERIFICA

Dipartimento di Epidemiologia
ASL RMF



Grade Profiles

- GRADE Profiles
 - Methadone maintenance treatment vs no treatment for opioids d
 - Use of opiate
 - Criminal behaviour
 - Mortality from RCTs
 - Mortality (Any cause) from observational studies
 - Mortality (Overdose) from observational studies
 - Retention in treatment

Outcome: Retention in treatment

Quality Assessment | Summary of findings | Other considerations | Quality of Evidence | Cochrane

Quality assessment

Outcome
Retention in treatment

Number of studies
3

How was the outcome assessed?
Objective

Design
Randomised trials

Limitations
No limitations

Consistency
No important inconsistency

Directness
No uncertainty

Other considerations
- NONE

Outcome: Retention in treatment

Quality Assessment | Summary of findings | Other considerations | Quality of Evidence | Cochrane

Outcome

Check for continuous outcome

Number of patients

Intervention		Comparison	
Methadone maintenance treatment		no treatment	
Number of patient	Total number of patients	Number of patients	Total number of patients
173	254	63	251
Percentage (68,1%)		Percentage (25,1%)	
Range: [] to []		Range: [] to []	

Effect

Relative (RR): 3.05

Absolute: 460 fewer / 1 000

95% CI (confidence limits): Low: 1.75 to High: 5.35

95% CI (confidence limits): Low: 270 more to High: 650 more

Length of follow up: 1 month- 2 years

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Effetto relativo

Fornisce la misura della forza dell'associazione tra il trattamento e l'outcome.

Può essere espresso in termini di Hazard Ratio, Rischio Relativo, Odds Ratio.

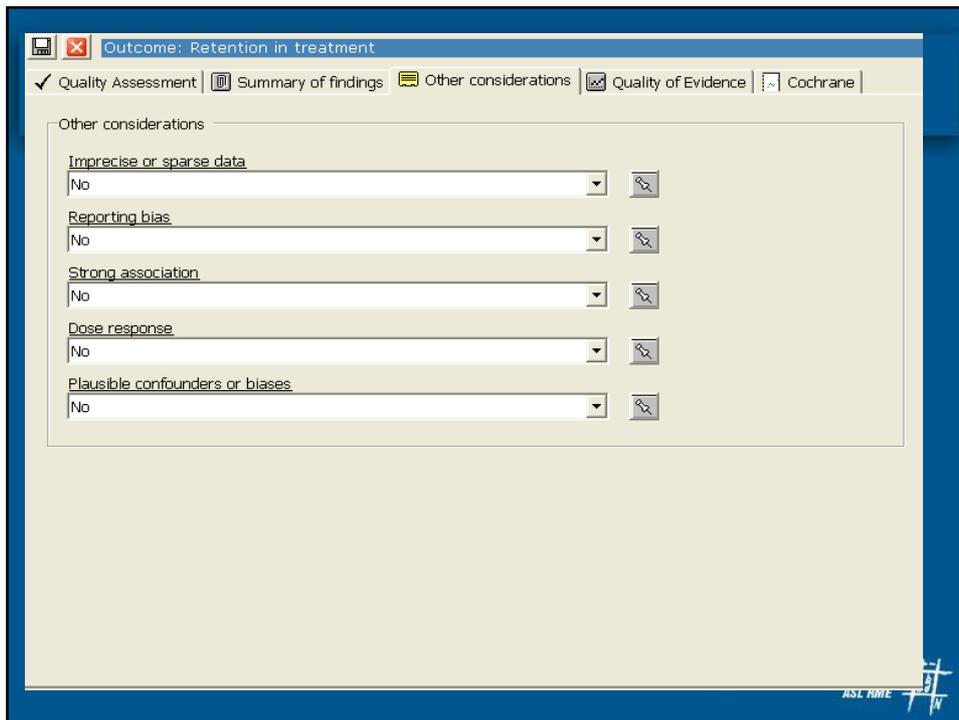
Effetto assoluto

Indica l'effetto assoluto dell'intervento; per esempio, quanti eventi in meno (o in più) si possono avere utilizzando un determinato trattamento nella popolazione considerata dal o dai trials.

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Quality assessment					
No of studies	Design	Limitations	Consistency	Directness	Other considerations
Use of opiate (Subjective Follow up: 1 month-2 years)					
3 ¹	Randomised trials ²	No limitations	No important inconsistency	No uncertainty	None
Criminal behaviour (Objective Follow up: 1 month-2 years)					
3 ¹	Randomised trials ²	No limitations	No important inconsistency	No uncertainty	None
Mortality from RCTs (Objective Follow up: 2-3 years)					
2 ⁴	Randomised trials ⁵	No limitations	No important inconsistency	No uncertainty	None
Mortality (Any cause) from observational studies (Objective Follow up: 2,5 years-21 years)					
5 ⁶	Observational studies ⁷	No limitations	No important inconsistency	No uncertainty	None
Mortality (Overdose) from observational studies (Objective Follow up: 2,5 years-12 years)					
5 ⁶	Observational studies ⁷	No limitations	Important inconsistency (-) ¹⁰	No uncertainty	None
Retention in treatment (Objective Follow up: 1 month- 2 years)					
3 ¹¹	Randomised trials ¹²	No limitations	No important inconsistency	No uncertainty	None
Footnotes:					
<ol style="list-style-type: none"> 1. 3 studies, outpatient setting, 2 conducted in USA and 1 in Sweden 2. 3 RCTs, 1 with adequate allocation concealment, 1 unclear, 1 inadequate 3. Random effect model 4. 2 RCTs, 1 conducted in USA and 1 in Sweden 5. 1 adequate and 1 unclear allocation concealment 6. 5 studies, outpatient setting, conducted in Italy, Australia, Sweden, Usa, Spain (1 each) 7. Quality of studies using Newcastle-Ottawa Scale: selection two studies rated 3 and three studies rated 2; comparability: one study rated 3, three rated 1 and one rated 0; outcome: two studies rated 2 and three rated 1 8. 5 studies, outpatient setting, 2 conducted in the Netherlands and one each in Italy, USA and Spain 9. Quality of studies using Newcastle-Ottawa Scale: selection: four studies rated 3 and one rated 2; comparability: two studies rated 2 and three rated 1; outcome: one study rated 2 and four rated 1 10. High statistical heterogeneity $p=0.00001$, but all consistent results 11. 3 studies, outpatient setting, conducted in Hong Kong, Thailand and Usa (one each) 12. 3 RCTs, all with unclear allocation concealment 					



Imprecise or sparse data

Criminal activity 1 (Objective¹⁵ Follow up: 12 months⁴)

1 ¹⁴	Randomised trials	No limitations ¹⁶	No important inconsistency	Major uncertainty (-2) ⁶	Imprecise or sparse data (-1) ¹⁷
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17. Only 1 study, few patients

Per decidere se i dati sono "imprecisi o sparsi":

- Un solo studio, con pochi partecipanti, un ampio intervallo di confidenza.

Reporting bias

The screenshot shows a software interface with a dropdown menu titled 'Reporting bias'. The menu is open, showing four options: 'No', 'No', 'High probability of reporting bias (- 1)', and 'No'. The first 'No' option is highlighted in blue. To the right of the dropdown are two small icons, one above and one below the menu.

Qualora vi sia:

- una selezione nella pubblicazione degli studi (**publication bias**)
- una pubblicazione selettiva degli outcome (**outcome reporting bias**)

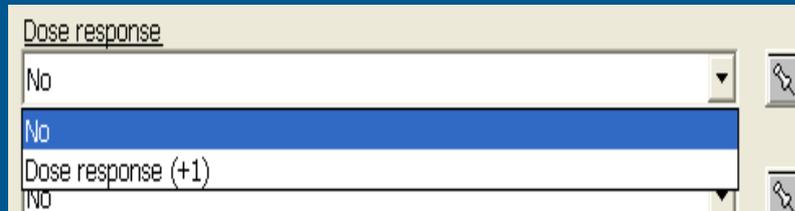
Strong association

The screenshot shows a software interface with a dropdown menu titled 'Strong association'. The menu is open, showing four options: 'No', 'No', 'Strong association (+1)', and 'Very strong association (+2)'. The first 'No' option is highlighted in blue. To the right of the dropdown are two small icons, one above and one below the menu.

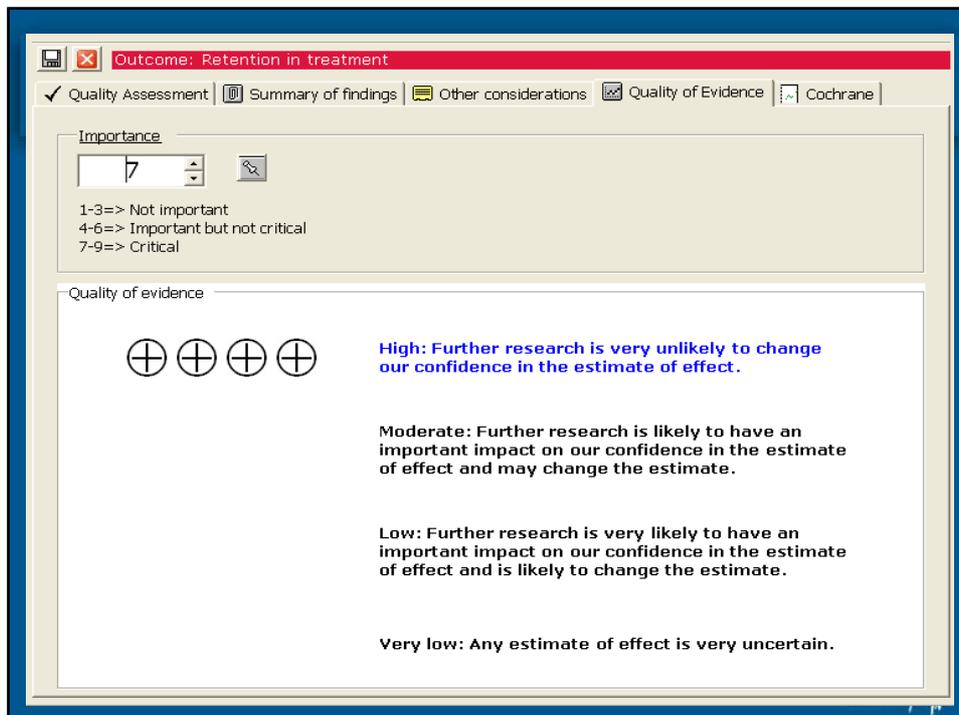
Un RR tra 2 e 5 (riduzione dal 50 all'80%) va considerato **forte associazione**

Un RR sopra 5 (riduzione > 80%) come **associazione molto forte**

Dose response



La presenza di una relazione dose-risposta può **migliorare** la qualità dell'evidenza



Outcome: Retention in treatment

Quality Assessment | Summary of findings | Other considerations | Quality of Evidence | Cochrane

Importance

7

1-3=> Not important
4-6=> Important but not critical
7-9=> Critical

Quality of evidence

⊕ ⊕ ⊕ ⊕

High: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low: Any estimate of effect is very uncertain.

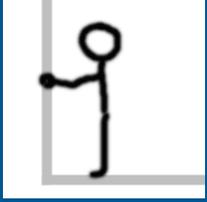
Quality assessment						Summary of findings					
No. studies	Design	Limitations	Consistency	Directness	Other considerations	No of patients		Effect		Quality	Importance
						Methadone maintenance treatment	No treatment	Relative risk (RR) (95% CI)	Absolute risk (AR) (95% CI)		
Use of opiates (subjective follow up: 1 month–2 years)											
3	Randomized trials	Some limitations ² (-1)	No important inconsistency	No uncertainty	None	28/104 (26.9%)	110/128 (87.3%)	RR 0.32 ¹ (0.23 to 0.44)	AR 630/1000 (830 less to 430 less)	⊕⊕⊕○ Moderate	7
Criminal behaviour (objective follow up: 1 month–2 years)											
3	Randomized trials	Some limitations ² (-1)	No important inconsistency	No uncertainty	Imprecise or sparse data (-1)	5/178 (2.8%)	18/185 (9.7%)	RR 0.39 ¹ (0.12 to 1.25)	AR 250/1000 (700 less to 19 more)	⊕⊕○○ Low	6
Mortality from randomized controlled trials (RCTs) (objective follow up: 2–3 years)											
3 ³	Randomized trials	No limitations	No important inconsistency	No uncertainty	Imprecise or sparse data (-2)	3/216 (1.4%)	7/219 (3.2%)	RR 0.48 ¹ (0.08 to 4.23)	AR 60/1000 (100 less to 90 more)	⊕⊕○○ Low	9
Mortality (any cause) from observational studies (objective follow up: 2.5 years–21 years)											
5 ⁴	Observational studies	No limitations	No important inconsistency	No uncertainty	None	257/19421 (1.3%)	1063/23814 (4.5%)	RR 0.37 (0.29 to 0.48)	AR 20/1000 (30 less to 10 less)	⊕⊕○○ Low	9
Mortality (overdose) from observational studies (objective follow up: 2.5 years–12 years)											
5 ⁵	Observational studies	No limitations	Inconsistent results between studies (-1)	No uncertainty	Extremely strong effect (+2)	70/37516 (0.2%)	416/32454 (1.3%)	RR 0.17 (0.05 to 0.63)	AR 10/1000 (20 less to 0.00)	⊕⊕○○ Moderate	9
Retention in treatment (objective follow up: 1 month–2 years)											
3 ³	Randomized trials	No limitations	No important inconsistency	No uncertainty	None	173/254 (68.1%)	63/251 (25.1%)	RR 3.05 ¹ (1.75 to 5.35)	AR 400/1000 (270 more to 650 more)	⊕⊕⊕⊕ High	7

Footnotes:

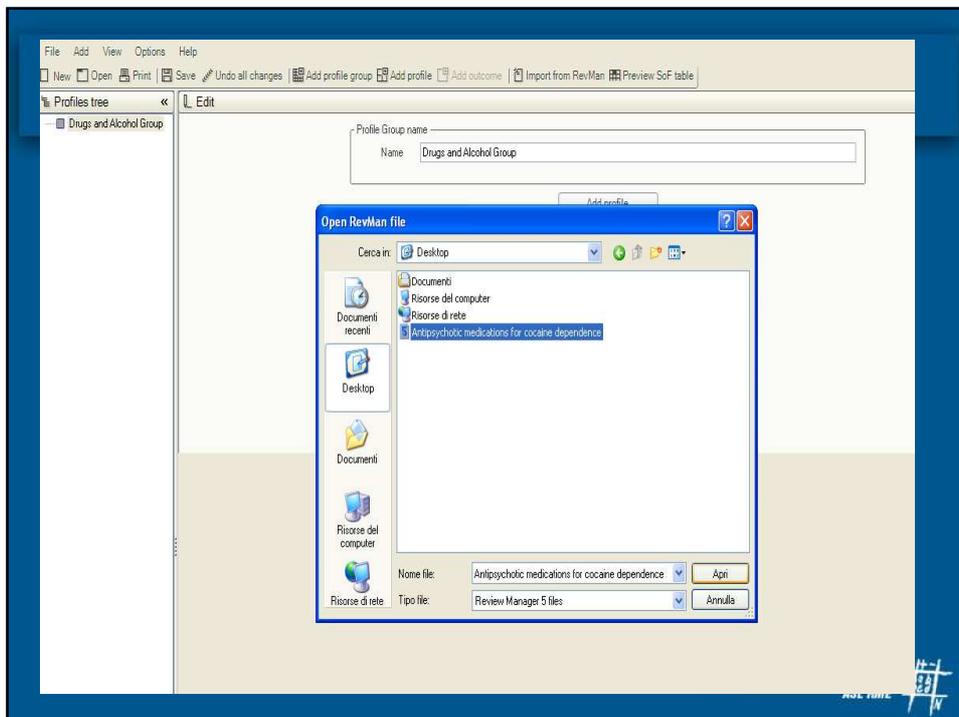
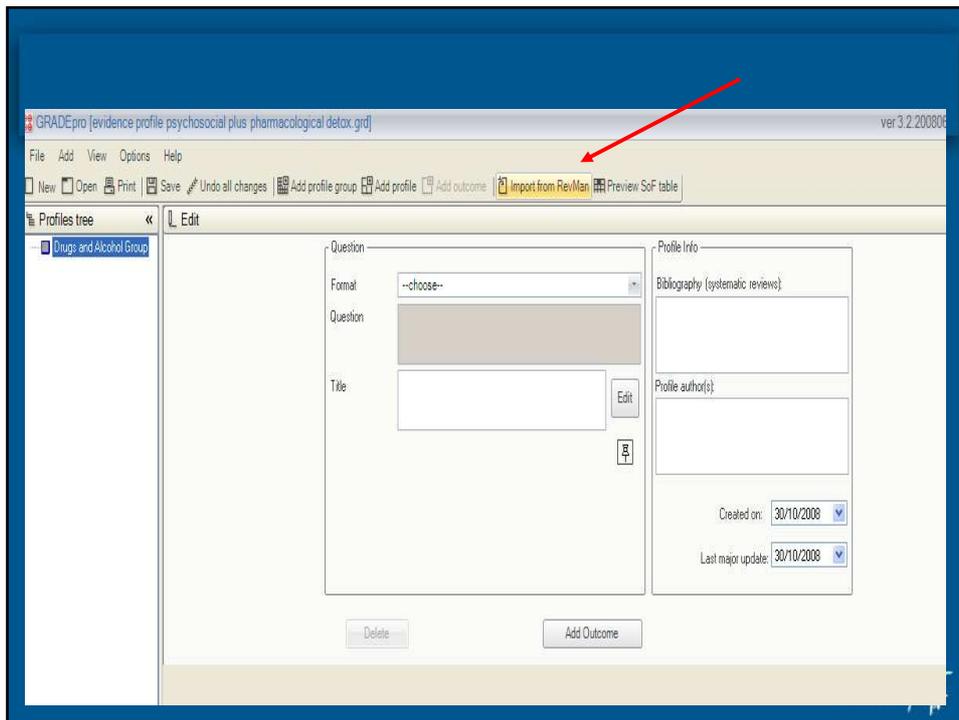
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12. 3 RCTs, all with unclear allocation concealment



[Redacted]







GRADEpro [evidence profile: psychosocial plus: pharmacological detox: gro] Ver 3.2.2008001

File Add View Options Help
 New Open Print Save Undo all changes Add profile group Add profile Add outcome Import from RevMan Preview SoF table

Profiles tree Edit

- Drugs and Alcohol Group
- Antipsychotic medications (1)
 - Any antipsychotic versus placebo
 - Dropouts
 - New Outcome
 - Risperidone versus Placebo
 - Olanzapine versus Placebo
 - Olanzapine versus Haloperidol

Outcome: Dropouts assessed with: []

Length of follow-up: [0] []

Number of participants: Intervention with event 44 total 106 41.5 %
 Control with event 56 total 102 54.9 %

Range of control group risks in individual studies: 6.7 % to 78.8 %

Control risk: Low 0 % Medium 20 % High 0 %

Estimate of the effect Relative: RR of 0.79 95% CI from 0.62 to 1.01
 Auto absolute effect calculation Absolute: 115 fewer per 1000 95% CI from 209 to 5

Delete Revert Go to Quality Assessment

Profile: Any antipsychotic versus placebo for cocaine dependence

Any antipsychotic versus placebo for cocaine dependence
 Patient or population: patients with cocaine dependence
 Settings:
 Intervention: Any antipsychotic versus placebo

Outcome	Assumed risk [Control]	Corresponding risk [Any antipsychotic versus placebo]	Relative effect (95% CI)	No of participants (studies)	Quality (GRADE)	Comments
Dropouts	200 per 1000	158 per 1000 (124 to 202)	RR 0.79 (0.62 to 1.01)	208 (5)		
[outcome not saved yet]	See comment	See comment	Not estimable	0 (0)	See comment	

Outcome: Dropouts dichotomous continuous pooled Importance: []

No of studies: 5
 Study design: -- choose -- Quality of evidence: []

Delete Revert Go to Summary of findings

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Any antipsychotic versus placebo for cocaine dependence
 Patient or population: patients with cocaine dependence
 Settings:
 Intervention: Any antipsychotic versus placebo

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