Abstract P1-276 Table 1  Respiratory diagnoses and prescriptions

<table>
<thead>
<tr>
<th>Drug misusers</th>
<th>Controls</th>
<th>Crude OR (95% CI)</th>
<th>Adjusted OR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Asthma</td>
<td>1590 (17.1%)</td>
<td>1009 (10.9%)*</td>
<td>1.695 (1.557 to 1.845)†</td>
</tr>
<tr>
<td>COPD (chronic obstructive pulmonary disease)</td>
<td>219 (2.4%)</td>
<td>74 (0.8%)*</td>
<td>3.007 (2.307 to 3.920)†</td>
</tr>
<tr>
<td>SABA (short acting beta agonist) prescribed</td>
<td>1520 (16.4%)</td>
<td>736 (7.9%)*</td>
<td>2.274 (2.071 to 2.486)†</td>
</tr>
<tr>
<td>LABA (long acting beta agonist) prescribed</td>
<td>92 (1%)</td>
<td>39 (0.4%)*</td>
<td>2.373 (1.630 to 3.454)†</td>
</tr>
<tr>
<td>ICS (inhaled corticosteroid) prescribed</td>
<td>987 (10.6%)</td>
<td>702 (7.6%)*</td>
<td>1.454 (1.314 to 1.609)†</td>
</tr>
</tbody>
</table>

* p<0.0001, † p<0.001 binary logistic regression. 诊断 ever recorded. § Prescriptions in 2008.

Conclusion These data suggest drug misusers have a significantly higher prevalence of respiratory diseases and are prescribed significantly more respiratory medications than matched controls. This exploratory study has set the scene for future work to explore possible reasons for this association.
decline from 23.9 % in 1998 to 21.8 % in 2009, mortality rates deserve our worries.

Conclusions Overall, the results point out to a declining co-infection trend. However treatment outcomes are quite worrisome; the cure of co-infected patients, being lower than the non co-infected, demonstrate the need to create priorities and adequate strategies for this population, mainly regarding the follow-up of these cases.

**P1-280** ANALYSIS OF TUBERCULOSIS RESISTANCE IN SAO PAULO STATE (SPS), BRAZIL FROM 2007 TO 2008

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**Introduction** Tuberculosis (TB) remains a serious public health problem. The most serious aspect of this problem has been the appearance of TB drug-resistance (TBDR), mainly the multidrug-resistance (TBMDR). One in five TB cases in Brazil occurs in Sao Paulo State, which has about 19,000 cases per year. Since 1980, TB treatment in Brazil has been with rifampin, isoniazid and pirazinamid. Thus, TB-DR monitoring is essential to assess its trend, in order to better assist the TB Control Program (TCP).

**Objective** To analyse TBDR in Sao Paulo State during the 2007–2008 period.

**Methodology** Using ‘population proportionate cluster sampling’ we selected 30 randomised clusters to design the resistance survey. During the 2007–2008 period, all respiratory symptomatic patients of these clusters answered the questionnaire and were submitted to AFB culture and sensitivity tests (ST).

**Results** From the 1746 patients interviewed, 30.3% were female and 69.7% male, the mean age was 37 years, 20.0% of the patients were HIV-positive and 821 were submitted to culture and ST. The primary resistance to isoniazid and rifampin was 5.8% and 1.5%, respectively. MDR-TB was 1.1% and any-resistance was 9.8%. The acquired-resistance was as follows: isoniazid, 20.4%; rifampin 12.2%; MDR-TB 12.2% and any-resistance 22.4%. The treatment outcomes of preeclampsia (RR = 1.03; 95% CI 0.95 to 1.11), small-for-gestational-age infants (RR = 0.95; 95% CI 0.66 to 1.05) and any baby death (RR = 1.02; 95% CI 0.87 to 1.20). Side-effects (abdominal pain, itching, eczema, vomiting, diarrhoea, headache, constipation, malaise, decreased vision, skin rash and chest pain) occurred more frequently among the women who took antioxidants than among those who took placebo (RR = 1.58; 95% CI 1.11 to 2.24).

**Conclusion** The evidence does not support the use of antioxidants during pregnancy. Not only are its benefits unclear, but also adverse effects occurred more frequently with its use.

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**P1-281** ANTI-OXIDANTS FOR PREVENTING PREECLAMPSIA: A SYSTEMATIC REVIEW

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**Objective** To investigate the effectiveness of antioxidants for preventing preeclampsia and other maternal and fetal complications among pregnant women with low, moderate or high risk of preeclampsia.

**Methods** We searched Medline, Embase, Cochrane Central, mRCT, CRD, ISI Web of Science, Lilacs, Scielo and Scopus databases, without language restriction or limits on date of publication. Randomised controlled clinical trials evaluating the use of antioxidants vs placebo or a group without antioxidants were considered eligible.

**Results** A total of 1120 articles were located, and 16 randomised clinical trials were analysed (20 808 women). A meta-analysis did not show any statistically significant difference between women who received an antioxidant (vitamin C, vitamin E, lycopene, selenium, red palm oil) and women who received placebo, for the outcomes of preeclampsia (RR = 0.92; 95% CI 0.80 to 1.06), severe preeclampsia (RR = 1.03; 95% CI 0.87 to 1.22), preterm birth (RR = 1.03; 95% CI 0.95 to 1.11), small-for-gestational-age infants (RR = 0.95; 95% CI 0.66 to 1.05) and any baby death (RR = 1.02; 95% CI 0.87 to 1.20). Side-effects (abdominal pain, itching, eczema, vomiting, diarrhoea, headache, constipation, malaise, decreased vision, skin rash and chest pain) occurred more frequently among the women who took antioxidants than among those who took placebo (RR = 1.58; 95% CI 1.11 to 2.24).

**Conclusion** The evidence does not support the use of antioxidants during pregnancy. Not only are its benefits unclear, but also adverse effects occurred more frequently with its use.

**Funding** MS/SCTIE/DECIT, via CNPq (Edital 67/2009).

**P1-282** SUBLINGUAL MISOPROSTOL FOR PREVENTING POSTPARTUM HAEMORRHAGE: A SYSTEMATIC REVIEW

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**Objective** To assess the efficacy and safety of sublingual misoprostol for preventing postpartum haemorrhage.

**Method** MEDLINE, Embase, CRD, CENTRAL, mRCT, LILACS, SciELO, ProQuest and ISI Web of Knowledge databases were searched. There were no language, accessibility, or publication date restrictions. Randomised clinical trials of sublingual misoprostol in comparison with placebo or other uterotonics were eligible. The primary outcome was postpartum haemorrhage (blood loss ≥500 ml). Other outcomes were considered (see below). The heterogeneity was evaluated and, when possible, the data grouped into a meta-analysis using a random-effects model.

**Results** Of 682 references identified, only 15 were included in the analysis (5109 patients). Most of the studies were of low methodologic quality. Sublingual misoprostol, at any dose, was not more effective for reducing postpartum haemorrhage in comparison with standard treatment. Sublingual misoprostol, however, is effective for reducing haemorrhage greater than 1000 ml [600 mcg vs placebo; RR = 0.66 (95% CI 0.45 to 0.98)] as well as the use of additional uterotonics [600 mcg vs methylergometrine; RR = 0.04 (0.00 to 0.72)] and the duration of the third stage of delivery [50 mcg vs placebo; RR = 0.95 (95% CI 0.87 to 1.03)]. The drug presented a worse safety profile, causing tremors and fever, especially at higher doses.

**Conclusion** Sublingual misoprostol aimed at preventing postpartum haemorrhage presents no benefits compared with the standard uterotonics. Its use should be restricted to clinics with adequate medical support, as a complement to other uterotonics.

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