Extrapyramidal motor side-effects of first- and second-generation antipsychotic drugs

Michael J. Peluso, Shôn W. Lewis, Thomas R. E. Barnes and Peter B. Jones

Background
Second-generation antipsychotics have been thought to cause fewer extrapyramidal side-effects (EPS) than first-generation antipsychotics, but recent pragmatic trials have indicated equivalence.

Aims
To determine whether second-generation antipsychotics had better outcomes in terms of EPS than first-generation drugs.

Method
We conducted an intention-to-treat, secondary analysis of data from an earlier randomised controlled trial (n = 227). A clinically significant difference was defined as double or half the symptoms in groups prescribed first- vs. second-generation antipsychotics, represented by odds ratios greater than 2.0 (indicating advantage for first-generation drugs) or less than 0.5 (indicating advantage for the newer drugs). We also examined EPS in terms of symptoms emergent at 12 weeks and 52 weeks, and symptoms that had resolved at these time points.

Results
At baseline those randomised to the first-generation antipsychotic group (n = 118) had similar EPS to the second-generation group (n = 109). Indications of resolved Parkinsonism (OR = 0.5) and akathisia (OR = 0.4) and increased tardive dyskinesia (OR = 2.2) in the second-generation drug group at 12 weeks were not statistically significant and the effects were not present by 52 weeks. Patients in the second-generation group were dramatically (30-fold) less likely to be prescribed adjunctive anticholinergic medication, despite equivalence in terms of EPS.

Conclusions
The expected improvement in EPS profiles for participants randomised to second-generation drugs was not found; the prognosis over 1 year of those in the first-generation arm was no worse in these terms. The place of careful prescription of first-generation drugs in contemporary practice remains to be defined, potentially improving clinical effectiveness and avoiding life-shortening metabolic disturbances in some patients currently treated with the narrow range of second-generation antipsychotics used in routine practice. This has educational implications because a generation of psychiatrists now has little or no experience with first-generation antipsychotic prescription.

Declaration of interest
In the past 3 years S.W.L. has received advisory board fees from Janssen-Cilag and speaker fees from AstraZeneca; T.R.E.B. has acted as a speaker at an event sponsored by Lilly; P.B.J. declares membership of a scientific advisory board for Roche, and has received research support from GlaxoSmithKline and a speaker fee from Lilly.

Antipsychotic medication has been the mainstay of schizophrenia treatment for the past 50 years. The first-generation antipsychotics, introduced in the mid-20th century, were unevenly effective in relieving the symptoms of schizophrenia, often at the expense of extrapyramidal side-effects (EPS) such as acute dystonia, akathisia, Parkinsonism and tardive dyskinesia. The development of second-generation antipsychotic drugs and their promotion by the pharmaceutical industry were predicated on indications that these medications would have a milder EPS profile and therefore greater tolerability than the earlier drugs.1-11 The results of recent large trials and meta-analyses have shown that there is no effectiveness or tolerability advantage.12-15 There is a view that the two generations of antipsychotics should preferably be seen as lying on a multidimensional continuum rather than as distinct classes, and that the heterogeneity and complexity of side-effects are masked by a spurious dichotomy.16,17 Nevertheless, there has been a dramatic increase in the prescription of second-generation drugs.18

The Cost Utility of the Latest Antipsychotics in Schizophrenia Study Band 1 (CUtLASS-1) was a pragmatic randomised controlled trial (RCT) that tested the hypothesis that the clinical and cost-effectiveness of second-generation antipsychotics would be superior in individuals whose antipsychotic treatment was being changed owing to an inadequate response or side-effects.12 The primary analysis of quality of life, symptoms and costs over 1 year refuted this hypothesis. In fact, the older drugs were associated with a trend towards better outcomes and lower costs, although the more conservative conclusion is one of broad equivalence between the two classes when prescribed in the context of a multicentre pragmatic trial. Second-generation drugs were not superior, even on measures of patient preference. One possible explanation for the relative lack of distinction between drug classes seen in CUtLASS-1 is that the second-generation antipsychotics were not associated with the expected relief from EPS. In the USA, the Clinical Antipsychotic Trials of Intervention Effectiveness (CATIE) study demonstrated, similarly, that there was no significant difference between second-generation antipsychotics when compared with perphenazine in terms of the emergence of EPS.19,20 It has been suggested in the analysis of CATIE and elsewhere that the differences between first- and second-generation drugs in early studies could have resulted from the use of haloperidol — often in relatively high dose — as the comparator.20,21 In fact, a systematic review published early in the second-generation drug epoch demonstrated no evidence that these drugs were more effective or better tolerated than the first generation, and attributed any perceived benefit to the dosage of the comparator drug.22 Furthermore, the doses of several second-generation antipsychotics used in clinical practice are higher than those used in the benchmarking studies sponsored by their manufacturers.

We report the results of a secondary analysis of data from the CUtLASS-1 study focusing on emergent and resolved EPS,
Method

The CUtLASS-l study was a pragmatic, multicentre, rater-masked RCT, conducted between July 1999 and January 2002 within 14 community psychiatry services affiliated with five medical schools in the English National Health Service. It was designed to test the effectiveness of antipsychotic medications in routine clinical practice. The 227 participants were randomised by means of a remote telephone service to receive either a first- or a second-generation antipsychotic (other than clozapine). Randomisation to a class of drugs allowed the managing physician to select a drug at randomisation and data were recoded according to the treatment to which they were allocated. Intention-to-treat (ITT) analysis was performed. Individuals were grouped depending on the treatment to which they were allocated at randomisation and data were recoded according to the operational criteria for EPS at baseline, 12 weeks and 52 weeks. To test whether emergent and relieved EPS differed between the two treatment groups, data were transformed into binary categories and presented in contingency tables from which chi-squared statistics and odds ratios were calculated; P values and 95% confidence limits were used to determine statistical significance.

It is common practice for clinicians to prescribe anticholinergic adjuncts to patients in response to EPS, particularly Parkinsonism, and sometimes in anticipation of such problems. To distinguish between patients receiving anticholinergic adjuncts in each study arm, the sample was stratified according to whether adjunctive medication was prescribed, effectively creating four treatment groups:

(a) first-generation antipsychotic alone;
(b) second-generation antipsychotic alone;
(c) first-generation antipsychotic with anticholinergic adjunct;
(d) second-generation antipsychotic with anticholinergic adjunct.

Procyclidine or trihexyphenidyl hydrochloride were the only anticholinergic drugs prescribed in this sample. After stratification by adjunct, the above analyses were repeated for the Parkinsonism outcome at 12 weeks and 52 weeks, as this is the condition that the adjuncts are most commonly used to prevent or treat. Comparisons were made between subgroups (a), (b) and (c).

Outcome measures

The main outcome measure for the primary analysis was the Quality of Life Scale (QLS) score, as previously reported. Secondary measures relevant to this analysis were the Barnes Akathisia Rating Scale (BARS) for akathisia, the Simpson–Angus Scale (SAS) for Parkinsonism and the Abnormal Involuntary Movement Scale (AIMS) for tardive dyskinesia. Side-effects were considered to be present at each time point according to the following operational criteria: for akathisia, when participants scored 2 or more on the global akathisia item of the BARS; for Parkinsonism, when participants had a total score of 3 or more on the SAS; and for tardive dyskinesia, when participants had one score of 3 or two scores of 2 on AIMS items 1–7 covering observed movements. An ‘emergent’ side-effect was defined as one that was not present at baseline but was noted at follow-up; a ‘resolved’ side-effect was present at baseline but absent at follow-up.

Results

Table 1 lists the antipsychotic drugs prescribed to patients randomised into first- or second-generation treatment groups and the doses at the end of the study, all of which are within conventional limits. The most common first-generation drugs chosen were sulpiride and trifluoperazine; haloperidol was a relatively uncommon choice. The most commonly prescribed second-generation drugs were olanzapine, quetiapine and risperidone.

Emergent side-effects

Table 2 describes the two treatment groups according to EPS at 12 weeks and 52 weeks, stratified into EPS that were emergent or resolved. There was no statistically significant difference between the groups in terms of emergent Parkinsonism, akathisia or tardive dyskinesia at either assessment point. Potentially clinically relevant differences in akathisia and Parkinsonism at 12 weeks in the second-generation group, both with odds ratios of 0.5 or less,
Table 1  Antipsychotic drugs prescribed at baseline in the two treatment arms

<table>
<thead>
<tr>
<th>Drug</th>
<th>Dose at end of study, mg/day at baseline</th>
<th>Patients prescribed drug at baseline</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean (range)</td>
<td>n</td>
</tr>
<tr>
<td>First-generation antipsychotic group (n = 118)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chlorpromazine</td>
<td>250 (200-300)</td>
<td>8</td>
</tr>
<tr>
<td>Droperidol</td>
<td>NA</td>
<td>1</td>
</tr>
<tr>
<td>Flupentol</td>
<td>4 (2-6)</td>
<td>1</td>
</tr>
<tr>
<td>Flupentol decanoate</td>
<td>142 2/52 (40-452-250 1/52)</td>
<td>2</td>
</tr>
<tr>
<td>Fluphenazine decanoate</td>
<td>50 2/52*</td>
<td>3</td>
</tr>
<tr>
<td>Haloperidol</td>
<td>22.5 (20-29)</td>
<td>7</td>
</tr>
<tr>
<td>Haloperidol decanoate</td>
<td>NA</td>
<td>2</td>
</tr>
<tr>
<td>Loxapine</td>
<td>NA</td>
<td>3</td>
</tr>
<tr>
<td>Pipotiazine palmitate</td>
<td>50 2/52</td>
<td>2</td>
</tr>
<tr>
<td>Sulpiride</td>
<td>813 (200-2400)</td>
<td>58</td>
</tr>
<tr>
<td>Thioridazine</td>
<td>NA</td>
<td>1</td>
</tr>
<tr>
<td>Trifluoperazine</td>
<td>15 (6-30)</td>
<td>21</td>
</tr>
<tr>
<td>Zuclopenthixol decanoate</td>
<td>358 2/52 (150 2/52-750 2/52)</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Second-generation antipsychotic group (n = 109)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Amisulpride</td>
<td>610 (200-1200)</td>
<td>13</td>
</tr>
<tr>
<td>Olanzapine</td>
<td>15 (5-30)</td>
<td>50</td>
</tr>
<tr>
<td>Quetiapine</td>
<td>450 (200-750)</td>
<td>23</td>
</tr>
<tr>
<td>Risperidone</td>
<td>5 (2-10)</td>
<td>22</td>
</tr>
</tbody>
</table>

a. Two data points are missing.
b. Equivalent dosing across all participants.

Table 2 Extrapyramidal side-effects in the first- and second-generation antipsychotic groups at baseline and at 12 weeks and 52 weeks follow-up, stratified by emergent and relieved symptoms at the two follow-up points

<table>
<thead>
<tr>
<th>Extrapyramidal side-effects</th>
<th>FGA group (n = 118)</th>
<th>SGA group (n = 109)</th>
<th>( \chi^2 )</th>
<th>( P )</th>
<th>SGA v. FGA OR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baseline symptoms</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Parkinsonism</td>
<td>61 (53)</td>
<td>57 (53)</td>
<td>0.0</td>
<td>0.93</td>
<td>1.0 (0.6-1.6)</td>
</tr>
<tr>
<td>Tardive dyskinesis</td>
<td>18 (15)</td>
<td>13 (12)</td>
<td>0.3</td>
<td>0.59</td>
<td>1.3 (0.6-2.9)</td>
</tr>
<tr>
<td>Akathisia</td>
<td>27 (23)</td>
<td>38 (33)</td>
<td>3.4</td>
<td>0.06</td>
<td>0.6 (0.3-1.0)</td>
</tr>
<tr>
<td>Emergent symptoms(^c)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12 weeks follow-up Parkinsonism</td>
<td>12 (11)</td>
<td>6 (6)</td>
<td>1.5</td>
<td>0.22</td>
<td>0.5 (0.2-1.5)</td>
</tr>
<tr>
<td>Tardive dyskinesis</td>
<td>4 (4)</td>
<td>7 (8)</td>
<td>1.6</td>
<td>0.21</td>
<td>2.2 (0.6-7.8)</td>
</tr>
<tr>
<td>Akathisia</td>
<td>8 (8)</td>
<td>4 (5)</td>
<td>1.8</td>
<td>0.18</td>
<td>0.4 (0.1-1.6)</td>
</tr>
<tr>
<td>52 weeks follow-up Parkinsonism</td>
<td>8 (8)</td>
<td>6 (6)</td>
<td>0.1</td>
<td>0.75</td>
<td>0.8 (0.3-2.5)</td>
</tr>
<tr>
<td>Tardive dyskinesis</td>
<td>8 (8)</td>
<td>7 (8)</td>
<td>0.0</td>
<td>0.97</td>
<td>1.0 (0.4-2.9)</td>
</tr>
<tr>
<td>Akathisia</td>
<td>5 (5)</td>
<td>4 (5)</td>
<td>0.0</td>
<td>0.89</td>
<td>0.9 (0.2-3.5)</td>
</tr>
<tr>
<td>Relieved symptoms(^d)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12 weeks follow-up Parkinsonism</td>
<td>14 (13)</td>
<td>12 (13)</td>
<td>0.0</td>
<td>0.95</td>
<td>1.0 (0.4-2.2)</td>
</tr>
<tr>
<td>Tardive dyskinesis</td>
<td>8 (7)</td>
<td>7 (6)</td>
<td>0.0</td>
<td>0.87</td>
<td>0.9 (0.3-2.6)</td>
</tr>
<tr>
<td>Akathisia</td>
<td>13 (11)</td>
<td>16 (15)</td>
<td>0.8</td>
<td>0.36</td>
<td>1.4 (0.7-3.2)</td>
</tr>
<tr>
<td>52 weeks follow-up Parkinsonism</td>
<td>21 (20)</td>
<td>15 (17)</td>
<td>0.4</td>
<td>0.51</td>
<td>0.8 (0.4-1.6)</td>
</tr>
<tr>
<td>Tardive dyskinesis</td>
<td>8 (7)</td>
<td>9 (9)</td>
<td>0.2</td>
<td>0.64</td>
<td>1.3 (0.5-3.4)</td>
</tr>
<tr>
<td>Akathisia</td>
<td>16 (14)</td>
<td>20 (20)</td>
<td>1.2</td>
<td>0.26</td>
<td>1.5 (0.7-3.1)</td>
</tr>
</tbody>
</table>

FGA: first-generation antipsychotic; SGA: second-generation antipsychotic.
a. Minor discrepancies in column percentages due to missing data; these percentages could theoretically exceed 100% if multiple extrapyramidal side-effects in some participants.
b. Baseline symptoms present at baseline but absent at follow-up.
c. Symptoms occurring in those free from symptom at baseline.
d. Symptoms present at baseline but absent at follow-up.

did not reach statistical significance and were no longer present at the 52-week follow-up, indicating a clinically significant increase in the development of tardive dyskinesia in the same group at 12 weeks (OR = 2.2, 95% CI 0.6-7.8) was similarly unconfirmed at conventional levels of statistical significance, and had disappeared by 52 weeks (OR = 1.0, 95% CI 0.4-2.9). These results suggest, overall, a null effect at 1-year follow-up.

There was no statistically significant difference between the treatment groups in terms of emergent Parkinsonism, akathisia or tardive dyskinesia at either follow-up point (Table 2). None of these effects achieved the a priori criteria for a clinically relevant effect in terms of symptom relief, suggesting that there was no clinically meaningful difference between the groups that was hidden by type 2 statistical error.
Use of anticholinergic adjuncts

In the first-generation antipsychotic group, 26 patients (22%) were prescribed an anticholinergic adjunct at baseline, a single patient (1%) in the second-generation group; this 30-fold increase in odds was highly statistically significant (P<0.001), despite the equivalence of EPS at this time point. Table 3 shows the results of an analysis of emergent Parkinsonism stratified by prescription of anticholinergic adjunct in the study population. No effect reached statistical significance but the trends were as follows. At 12 weeks, equivalence of EPS at this time point. Table 3 shows the results of a first-generation antipsychotic plus anticholinergic medication, an analysis of emergent Parkinsonism stratified by prescription of criteria for a clinically relevant effect. At 52 weeks, there was no patient (1%) in the second-generation group; this 30-fold increase in odds was highly statistically significant (P<0.001), despite the equivalence of EPS at this time point. Table 3 shows the results of a first-generation antipsychotic plus anticholinergic medication, an analysis of emergent Parkinsonism stratified by prescription of criteria for a clinically relevant effect. At 52 weeks, there was no clinically significant difference between first- and second-generation antipsychotics in terms of relief from baseline EPS at either 12-week or 52-week follow-up. Second-generation drugs appeared to be no more successful than the older ones in providing relief from these side-effects. This is surprising in the context of the common belief that first-generation antipsychotics exacerbate such problems, but nonetheless is in line with the CATIE results.

Use of anticholinergic adjuncts

We report the results of a secondary analysis of EPS in the CATLASS-1 trial. Owing to the statistical power constraints in such an analysis, we framed our results in the context of clinically important effects, defined as an odds ratio of 2.0 or 0.5, a double or half risk of EPS between the two study groups. Statistical power was limited but this approach allows us to conclude that there were few clinically relevant differences missed due to type 2 errors. The results were essentially null. The more frequent prescription of anticholinergic agents in the first-generation drug group despite equivalent EPS at baseline almost certainly represents clinical expectation of greater EPS in this arm. Overall, these results are in accord with those from other studies.12-15,19,20

Drug choice

Many RCTs showing second-generation antipsychotics to have a lower risk of EPS than a first-generation drug used haloperidol as the comparator. Haloperidol has a relatively high EPS liability and is often prescribed in a high dose that may be above the optimum dose for the study sample. The findings from the CATLASS-1 study indicate that, in this pragmatic trial designed to emulate real physician prescribing behaviour, haloperidol was an uncommon choice for first-generation antipsychotic treatment (Table 1). This could contribute to the results suggesting that there are, generally, few clinically significant differences in the EPS profiles of first- and second-generation antipsychotics when those drugs are prescribed with a flexible approach and due care.

Emergent side-effects

The results from the analysis of emergent EPS showed that at 1-year follow-up there was no clinically significant difference between the two drug classes. Although there was a clinically significant decrease in Parkinsonism and akathisia for the second-generation antipsychotic group at 12 weeks (ORs 0.5 and 0.4 respectively), this effect was diminished at the 52-week follow-up point, suggesting that the benefit was not long-lasting. It does, however, remain possible that the clinically significant decrease in EPS at 12 weeks was real and that the null effect at 52 weeks was due to increased class switching that is not reflected in the ITT analysis but has been previously reported.21-23

Despite the decreases in Parkinsonism and akathisia at 12 weeks, tardive dyskinesia was twice as common at this point in the second-generation group (OR = 2.2) but was not statistically significant and this potentially clinically relevant effect also disappeared by 52 weeks. One possible interpretation of this result is that tardive dyskinesia is temporarily exacerbated by withdrawal of dopamine-2 receptor blockade, reflecting a change in the neurotransmitter milieu resulting from the switch in drug class. In addition, the degree of tardive dyskinesia has been shown to worsen with adjunctive anticholinergic medication,29 and to improve with its discontinuation.30 Therefore, the high intrinsic anticholinergic activity of some second-generation drugs such as olanzapine may have contributed to this effect.31

Relieved side-effects

There was no clinically significant difference between first- and second-generation antipsychotic drugs in terms of relief from baseline EPS at either 12-week or 52-week follow-up. Second-generation drugs appeared to be no more successful than the older ones in providing relief from these side-effects. This is surprising in the context of the common belief that first-generation antipsychotics exacerbate such problems, but nonetheless is in line with the CATIE results.

Use of anticholinergic adjuncts

Anticholinergic adjuncts were more typically prescribed to prevent or mitigate EPS such as Parkinsonism in those receiving first-generation antipsychotics. However, the justification for the overwhelming difference in adjunct prescription between the two treatment arms is unclear, given the fact that there was no sustained, clinically relevant difference in EPS between the two groups and no difference at baseline. An anticholinergic adjunct was prescribed for just one patient taking a second-generation drug, despite the equivalence in EPS profiles between the classes in this study. One possible explanation for this finding is that clinicians are more likely to prescribe anticholinergic adjuncts on the basis of their expectations regarding the side-effect profile of the drug, especially given their likely assumption at the time that second-generation antipsychotics would have markedly lower liability to EPS.32 Clinicians prescribing a first-generation drug may have expected the development of EPS and/or had a lower

<table>
<thead>
<tr>
<th>Treatment subgroup</th>
<th>Parkinsonism &lt; 1.0 (n)</th>
<th>OR (95% CI)</th>
<th>P</th>
<th>OR (95% CI)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 weeks follow-up</td>
<td>FGAs</td>
<td>11 (13)</td>
<td></td>
<td>2.4 (0.4-13.3)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>SGA - adjunct</td>
<td>6 (6)</td>
<td></td>
<td>0.7 (0.2-5.0)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>SGA + adjunct</td>
<td>1 (4)</td>
<td></td>
<td>0.7 (0.2-2.3)</td>
<td></td>
</tr>
<tr>
<td>52 weeks follow-up</td>
<td>FGAs</td>
<td>7 (9)</td>
<td></td>
<td>0.4 (0.2-1.3)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>SGA + adjunct</td>
<td>0 (0)</td>
<td></td>
<td>0 (0)</td>
<td></td>
</tr>
</tbody>
</table>

FGA, first-generation antipsychotic; SGA, second-generation antipsychotic.

Table 3 Emergent Parkinsonism at 12 weeks and 52 weeks follow-up stratified by treatment arm and prescription of anticholinergic adjunct.
threshold for the detection of these symptoms; thus, they would
have been more likely to prescribe an anticholinergic drug as an
adjunct in anticipation of or in response to EPS. Other studies
have shown that adherence to second-generation antipsychotic
treatment can be enhanced by the prescription of an anticholinergic
along with the antipsychotic, so it is notable that in this case they
were so rarely used. The clinical threshold for detection of EPS
can be higher than that applied when prescribing a first-generation antipsychotic where these symptoms are expected. This was true for psychotic symptoms, as clinicians switched patients randomised to first-generation antipsychotics at a lower score on the Positive and Negative Syndrome Scale than patients randomised to the newer drugs.23

The analysis of emergent Parkinsonism stratified by adjunctive
prescription showed a clinically significant difference when
comparing second-generation antipsychotics with first-generation antipsychotics alone at 12-week follow-up. However, there was no clinically significant difference in the comparison between second-generation antipsychotics and first-generation antipsychotics plus anticholinergic at this time point. This suggests that potential EPS from first-generation antipsychotics can be effectively managed with adjunctive anticholinergic medication. At 52 weeks there was no clinically significant difference between any of the three
groups. Anticholinergic prescription may have attendant problems, such as cognitive deficit and potential for misuse, but we note this was not reflected in the overall results of the CUtLASS-1 study.12

Limitations
Secondary analyses of trial data often face limitations in terms of
statistical power given that the original studies are designed
around the primary outcome of the original trial. Estimates of
power undertaken post hoc indicated that this was reasonable for
the clinically relevant effects that we defined. By pre-defining these
clinically significant odds ratios we were able to interpret our
results in the context of patient experience with these medications;
we do not think we have missed important effects due to type 2
error. An effort was also made to minimise the number of
statistical tests carried out in this analysis to control for type 1
errors. For instance, the use of anticholinergic drugs by clinicians
in response to Parkinsonism is based on the strongest evidence, so
we restricted statistical tests to this side-effect. We did not stratify
our analyses according to the reason for referral to the trial for
similar reasons of limited statistical power.

Masking of raters to treatment allocation is an important
source of potential bias. Considerable efforts were made to
maintain the masking, including physical separation of raters from
clinical teams, reminders to patients not to divulge their
treatment, and technical aspects of the randomisation procedure
and study database. Known breaches were reported and affected
four participants in the first-generation group and two in the
second-generation group. Nevertheless, it is possible that subtle
indications were apparent in more cases. If such bias was present,
however, we believe that it would most probably have operated
against the older drugs, for example EPS might be more likely
to be rated as present in participants in whom signs of treatment
with an anticholinergic agent were present. Thus, we consider this
potential bias an unlikely cause of the null results.

Implications
This analysis illuminates the relative side-effect profiles of first-
and second-generation antipsychotics in terms of EPS when used
in the context of a clinical trial. It suggests some misconceptions
prevalent among the participating clinicians at that time regarding
their expectation of motor side-effects; they were, in fact, able to
use the two classes of drugs with equivalence in EPS. This ITT
comparison shows that there was weak evidence (not statistically
significant) for few clinically significant differences in terms of
emergent or relieved EPS between the two classes of antipsychotics
at 12 weeks, and none at 52 weeks. One implication is that
judicious prescription of adjunctive anticholinergic agents to
manage EPS when prescribing first-generation antipsychotics can
result in an EPS profile equivalent to second-generation drug
treatment.23 This analysis contributes to the existing literature on
EPS profiles of schizophrenia medications by demonstrating the
effects of these drugs in real clinical practice. However, further
work is necessary to determine definitively the treatment regimens
that will provide the greatest benefit while causing the fewest
adverse effects for people with schizophrenia and similar
disorders.

The results suggest that prescribers can rise to the challenge of
using both first- and second-generation compounds at doses that
result in levels in the domain between the dose-response curves
for beneficial and extrapyramidal effects. This dose-dependent
therapeutic domain differs not only between drugs but also
between patients. Although the introduction of second-generation
drugs may have improved prescribing through an increased
emphasis on monotherapy and vigilance for EPS, these benefits
can be obtained with first-generation drugs and may be better
achieved with the latter in some cases. Indeed, many patients
may not have been well served by the rapid restriction of the
number of antipsychotic drugs in common use, as the second-
generation drugs became the only antipsychotics used by most
clinicians.25

The emergence of obesity and metabolic abnormalities leading
to life-threatening increases in risk of cardiovascular disease is a
serious complication of antipsychotic prescribing, with second-
generation drugs in particular being implicated.23 In advance of
new antipsychotic drugs with truly novel mechanisms of action
that may not have these side-effects, patients urgently need us
to reappraise the relative positions of the two generations of drugs
on our therapeutic palette, ideally with further randomised trials
to guide the use of a wider range of antipsychotic options. There
are educational implications of a return to the careful use of first-
generation drugs as a treatment option for some people, because a
generation of psychiatrists is now unfamiliar with their use.

1 Arvanitis LA, Miller BG. Multiple fixed doses of 'Seroquel' (quetiapine) In
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