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Recurrence of Hyperprolactinemia after Withdrawal of Dopamine Agonists: Systematic Review and Meta-Analysis

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Context: Dopamine agonists are the treatment of choice for prolactinomas and symptomatic idiopathic hyperprolactinemia. However, the optimal treatment strategy and treatment duration is not clear in all details.

Objective: The aim of the study was to assess the effect of dopamine agonist withdrawal in patients with idiopathic hyperprolactinemia and prolactinomas.

Data Sources: PubMed, the Cochrane Library, the Web of Science, and EMBASE were searched electronically. No restriction was made with respect to language.

Study Selection: Studies reporting the proportion of normoprolactinemic patients after withdrawal of dopamine agonist or studies in which this proportion could be calculated were eligible. Both observational studies and clinical trials were eligible. Nineteen studies were included in the meta-analysis, with a total of 743 patients.

Data Extraction: Data extraction was performed by two reviewers independently.

Data Synthesis: The pooled proportion of patients with persisting normoprolactinemia after dopamine agonist withdrawal was 21% in a random effects model [95% confidence interval (CI), 14–30%; I² 81%]. Stratified analysis showed higher proportions of treatment success in idiopathic hyperprolactinemia (32%; 95% CI, 5–80%), compared with both microprolactinomas (21%; 95% CI, 10–37%), and macroprolactinomas (16%; 95% CI, 6–36%). In a random effects meta-regression adjusting for cause of hyperprolactinemia, a longer treatment duration was associated with treatment success (P = 0.015), whereas the use of cabergoline showed a trend of effect (P = 0.07).

Conclusions: This meta-analysis showed that hyperprolactinemia will recur after dopamine agonist withdrawal in a considerable proportion of patients. The probability of treatment success was highest when cabergoline was used for at least 2 yr. (J Clin Endocrinol Metab 95: 43–51, 2010)

Dopamine agonists are the treatment of choice for prolactinomas and symptomatic idiopathic hyperprolactinemia. Whether dopamine agonist treatment should be lifelong has been a subject of debate (1). Since the introduction of dopamine agonists more than 30 yr ago, several studies have assessed the effect of dopamine agonist withdrawal on the recurrence of hyperprolactinemia. The debate has shifted from whether a dopamine agonist can be stopped to the determination of the optimal timing for withdrawal (2). However, the optimal treatment strat-
egy is not clear in all details, resulting in different treat-
ment policies (3). A recent study (4, 5) indicated that a
subgroup of hyperprolactinemic patients with a high like-
lihood of achieving remission can be identified on clinical
criteria. In 2006 the Pituitary Society published consensus
guidelines (6) that summarized these controversies.

To assess the effect of dopamine agonist withdrawal in
patients with idiopathic hyperprolactinemia and prolacti-
nomas in more detail, we performed a systematic review of
the literature. The primary aim was to estimate the pooled
proportion of patients with persistent normoprolactinemia
after withdrawal of dopamine agonists in a meta-analysis.
The second aim was to determine factors influencing the suc-
cess of treatment outcome in a sensitivity analysis.

Materials and Methods

Eligibility criteria

The main outcome of the present analysis was the proportion
of patients with persisting normoprolactinemia after withdrawal
of dopamine agonist treatment in idiopathic hyperprolactinemia and prolactinomas. Studies reporting the proportion of normo-
prolactinemic patients after withdrawal of dopamine agonist or studies in which this proportion could be calculated were eligible
for inclusion in this meta-analysis. The assessment of recurrence
of hyperprolactinemia was based only on prolactin levels, irre-
spective of clinical symptoms. Both observational studies and
clinical trials were eligible. There were no restrictions with re-
spect to the sort of dopamine agonist.

Studies were eligible for inclusion in this review if they ful-
filled the following criteria:
1. The normal reference values of prolactin had to be reported.
2. Duration of dopamine agonist treatment was at least 3
months, and during the treatment period normoprolactinemia
had to be attained.
3. Follow-up period for patients with persisting normopro-
lactinemia after treatment withdrawal was at least 6 months.
4. The maximum proportion of pretreatment with radio-
therapy in patients assessed for the effect of dopamine agonist
withdrawal was set at 20%. The reason for this constraint is
that the effect of radiotherapy on hyperprolactinemia can be
delayed for many years. Therefore, radiotherapy is a con-
founder in the assessment of the effect of dopamine agonist
withdrawal.
5. Variables as age, sex, type of dopamine agonist, and
treatment duration had to be reported. If only a subgroup of
a larger cohort was withdrawn from dopamine agonist treat-
ment, these parameters were extracted for this subgroup only.
If, however, these parameters were not reported for the sub-
group separately, the parameters of the total cohort were ex-
tracted as a proxy for the subgroup. The latter condition was
only permitted if the minimum percentage of patients in the
study cohort who attained normoprolactinemia during treat-
ment and subsequently stopped treatment was at least 75% of
the total study group. In this way, the variables of the total
cohort should be a reliable estimate of the variables of those
who attained normoprolactinemia during treatment and sub-
sequently stopped treatment.

6. There should be no (partial) duplication of cohorts. If,
nonetheless, partial duplication was present, the largest cohort
was included.

If the entire cohort did not fulfill the eligibility criteria, the
included cohort was restricted to eligible patients. However, this
was only possible in case the study provided data and outcomes
on individual patients or if data and outcomes were shown ac-
cording to subgroups; i.e., if possible, the cohort was restricted
to patients without radiotherapy, nonpregnant during follow-
up, and with a normalized prolactin before withdrawal of the
medication.

Search strategy

We searched the PubMed, the Cochrane Library, the Web of
Science, and EMBASE databases for publications in any lan-
guage examining the effect of withdrawal of dopamine agonists
on the recurrence of hyperprolactinemia. The search was re-
stricted by date of publication from 1970 onward because do-
pamine agonists were not available for the treatment of hyper-
prolactinemia before 1970. For details of the search strategy, see
the Appendix (published as supplemental data on The Endocrine

Searches were performed in July 2008. In addition, the re-
ferences of relevant articles were checked for additional articles.
Abstracts of meetings and unpublished results were not included
in the analysis. There was no restriction with respect to language.

Data review and data analysis

Initial selection of studies by title and abstract was carried out
by one reviewer (J.L.). These studies were retrieved for full as-
essment. This assessment and subsequent data extraction were
performed by two independent reviewers (J.L. and O.M.D.).
Disagreements were resolved by consensus. Whenever possible,
the study cohort was stratified by cause of hyperprolactinemia:
idiopathic hyperprolactinemia and hyperprolactinemia caused
by micro- and macroprolactinomas, respectively. The provided
reference ranges of the individual studies were used to determine
the presence of hyper and normoprolactinemia, despite the fact
that in a few studies the authors defined remission as mild hy-
perprolactinemia without symptoms. All prolactin levels in the
present meta-analysis were expressed as micrograms per liter.

The conversion factor for prolactin levels from milliunits per liter
to micrograms per liter used was 1:30 (5), because we were not
able to acquire all conversion factors for the various assays. For
assessment of recurrence of hyperprolactinemia, the unit used by
the authors was used, not the converted levels. For studies re-
porting outcomes for several time intervals, the last point in time
was chosen for data extraction. For determination of tumor re-
gression during dopamine agonist treatment, results from mag-
netic resonance imaging (MRI) and computed tomography (CT)
were taken into account, not from conventional x-rays. For all
studies, the number of pregnant patients during follow-up was
extracted from the article if possible.

The main outcome of the meta-analysis was the weighted
average of the proportion of patients with persisting normo-
prolactinemia after withdrawal of dopamine agonist therapy.

The individual studies were weighted according to the inverse
of the squared se. The F test was used to check for quantitative
heterogeneity (7). This measures the proportion of inconsis-
tency between the studies that cannot be explained by chance
alone.
We performed the analyses stratified by causes of hyperprolactinemia (idiopathic hyperprolactinemia, microprolactinomas and macroprolactinomas), the type of dopamine agonist, and treatment duration (up to and including 24 months vs. more than 24 months). Random effects meta-regression was performed to study the influence of treatment duration and dopamine agonist preparation on persisting normoprolactinemia.

Statistical analyses were done in Comprehensive Meta-Analysis (version 2.0; Biostat, Englewood, NJ) and Stata (version 10.0; Stata Corporation, College Station, TX).

Results

Literature search (Fig. 1)

The initial search in the databases resulted in a total of 968 articles (543 in PubMed, 29 in Cochrane Library, 84 in Web of Science, and 312 in EMBASE). Of these 968 studies, 754 were unique without duplications. We excluded 685 papers based on title and abstract. A total of 69 potentially relevant papers were retrieved for full assessment. Of these studies, 31 were excluded from further analysis because the studies did not contain original data on withdrawal of dopamine agonists in hyperprolactinemia. We were unable to obtain one study (8).

In 40 studies, a detailed assessment with respect to the eligibility criteria was performed. Twenty studies were excluded from further analysis because these did not meet one or more of the eligibility criteria (4, 9–28). Two studies partially described the same cohort (4, 5); from these, the study representing the extension was included (5). Consequently, a total of 19 studies were included in the present review (5, 29–46).

For some studies the following subgroups were not included: microprolactinomas treated with bromocriptine (38) and macroprolactinomas (31, 33, 36). Reasons for exclusions of these subgroups were the application of radiotherapy in a large proportion of the subgroup (31, 33, 36), and a duration of follow-up after withdrawal in the subgroup shorter than 3 months (38). In two studies, patients were assessed twice after treatment with two different dopamine agonists. To prevent multiplicity of patients in the meta-analysis from these two studies, only one of the assessed treatments was included. In one study with a crossover design, only the results for cabergoline, but not for treatment with quinaagolide, were included because during treatment with quinaagolide two patients were nonevaluable (35). In a second study, only the outcomes after the first treatment, i.e. quinaagolide, were included (32). Of all included studies, three studies were retrieved after inspection of the references of relevant literature.

Study characteristics

Details of the 19 included studies are summarized in Table 1. Studies on persisting normoprolactinemia after withdrawal of dopamine agonists were published between 1979 and 2007. In nine studies, patients were treated according to a prespecified protocol (5, 30, 32, 34, 35, 37, 38, 40, 41). The number of included patients per study ranged from 2 to 221. The total number of patients included in this meta-analysis was 743. There were stratified data available for a total of 49 patients with idiopathic hyperprolactinemia, 353 with microprolactinomas, and 159 with macroprolactinomas. Idiopathic hyperprolactinemia was defined by authors as an unexplained hyperprolactinemia in the presence of normal CT or MRI (5, 35, 38). In none of these three studies was macroprolactinemia explicitly ruled out. In three studies with a total of 182 patients, the patients could not be separated with respect to different etiology. In two studies, a considerable proportion of patients were pregnant during follow-up: 43% (42) and 30% (29). In one study, the data for analysis could be restricted to nonpregnant patients (30). In a few other studies one (5, 34, 40) or two (41) patients became pregnant during follow-up.

Meta-analysis (Table 2)

The proportion of patients with persistent normoprolactinemia after withdrawal of dopamine agonists ranged

<table>
<thead>
<tr>
<th>754 potentially relevant studies identified and screened</th>
</tr>
</thead>
<tbody>
<tr>
<td>685 papers excluded on the basis of title and abstract</td>
</tr>
<tr>
<td>69 potentially relevant papers retrieved for more detailed assessment</td>
</tr>
<tr>
<td>31 papers excluded because not relevant for research question 1 paper not possible to obtain</td>
</tr>
<tr>
<td>37 papers with data on withdrawal of dopamine agonists in hyperprolactinemia</td>
</tr>
<tr>
<td>3 papers included after check of references of articles</td>
</tr>
<tr>
<td>40 papers with data on withdrawal of dopamine agonists in hyperprolactinemia</td>
</tr>
<tr>
<td>21 papers excluded because not fulfilling the inclusion criteria</td>
</tr>
<tr>
<td>19 studies with data on withdrawal of dopamine agonists in hyperprolactinemia fulfilling the inclusion criteria</td>
</tr>
</tbody>
</table>

FIG. 1. Summary of study assessment and exclusion stages.
<table>
<thead>
<tr>
<th>First author, year of publication (Ref.)</th>
<th>Cause of hyperprolactinemia</th>
<th>Treatment and mean dose (mg/d)</th>
<th>No. of patients</th>
<th>Mean age (yr)</th>
<th>Female sex (%)</th>
<th>Mean treatment duration (months)</th>
<th>Pretreatment</th>
<th>Mean PRL before treatment (ng/liter)</th>
<th>Regression of tumor during treatment on MRI or CT</th>
<th>Persisting normoprolactinemia, n (%)</th>
<th>Mean follow-up in persisting normoprolactinemia (months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Colao, 2007 (5)</td>
<td>Idiopathic</td>
<td>CAB 0.5 mg</td>
<td>27</td>
<td>27</td>
<td>100</td>
<td>39</td>
<td>None</td>
<td>67.8</td>
<td>At least 50% reduction in tumor volume</td>
<td>2/27 (74.1%)</td>
<td>57</td>
</tr>
<tr>
<td></td>
<td>Micro</td>
<td>CAB 1.2 mg</td>
<td>115</td>
<td>32</td>
<td>89.6</td>
<td>43</td>
<td>None</td>
<td>157.2</td>
<td>At least 50% reduction in tumor volume</td>
<td>76/115 (66.1%)</td>
<td>47</td>
</tr>
<tr>
<td></td>
<td></td>
<td>CAB 1.2 mg</td>
<td>79</td>
<td>44</td>
<td>54.4</td>
<td>42</td>
<td>None</td>
<td>89.17</td>
<td></td>
<td>37/79 (46.8%)</td>
<td>44</td>
</tr>
<tr>
<td>Biswas, 2005 (29)</td>
<td>Micro</td>
<td>BRC (n = 22), range 2.5–10 mg; CAB (n = 67), range 0.5–3 mg</td>
<td>89</td>
<td>32.7</td>
<td>94.4</td>
<td>37.2</td>
<td>None</td>
<td>71.3</td>
<td>At least 50% reduction in tumor volume</td>
<td>23/89 (26%)</td>
<td>43.2</td>
</tr>
<tr>
<td>Passos, 2002 (42)</td>
<td>Total (n = 131), micro (n = 62), macro (n = 69)</td>
<td>CAB 6.9 mg</td>
<td>131</td>
<td>32.1</td>
<td>77.1</td>
<td>60</td>
<td>None</td>
<td>89.4</td>
<td></td>
<td>Total 27/131 (20.6%), micro 16/62 (25.8%), macro 11/69 (15.9%)</td>
<td>37</td>
</tr>
<tr>
<td>Di Sarno, 2000 (32)</td>
<td>Micro</td>
<td>QUI 0.121 mg</td>
<td>23</td>
<td>34.6</td>
<td>91.3</td>
<td>12</td>
<td>1/23 OP</td>
<td>155.6</td>
<td>5/23 (22%) at least 80% reduction</td>
<td>0</td>
<td>NA</td>
</tr>
<tr>
<td></td>
<td>Macro</td>
<td>QUI 0.257 mg</td>
<td>14</td>
<td>32.1</td>
<td>64.3</td>
<td>12</td>
<td>5/16 OP</td>
<td>92.1</td>
<td>4/16 (25%) at least 80% reduction</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>Cannavo, 1999 (30)</td>
<td>Micro</td>
<td>CAB 0.98 mg</td>
<td>18</td>
<td>30.8</td>
<td>87.0</td>
<td>24</td>
<td>None</td>
<td>193.8</td>
<td>11/23 (48%) total disappearance</td>
<td>4/11 (36%) total disappearance</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>Macro</td>
<td>CAB 1.77 mg</td>
<td>9</td>
<td>28.1</td>
<td>90.9</td>
<td>24</td>
<td>None</td>
<td>40.4</td>
<td></td>
<td>1/9 (11.1%)</td>
<td>12</td>
</tr>
<tr>
<td>Musatori, 1997 (41)</td>
<td>Micro</td>
<td>CAB 0.93 mg</td>
<td>25</td>
<td>Range 25–48</td>
<td>100</td>
<td>12</td>
<td>3/26 OP, 9/26 BRC, 9/26 DHEC</td>
<td>124.8</td>
<td>2/25 (8.3%)</td>
<td>2/25 (8.3%)</td>
<td>49</td>
</tr>
<tr>
<td>Giusti, 1994 (35)</td>
<td>Idiopathic 6, micro 5, empty sella 1</td>
<td>CAB 1.0 mg</td>
<td>11</td>
<td>30.2</td>
<td>100</td>
<td>3</td>
<td>OP because of macroadenoma (n = 1) 10/12 BR</td>
<td>80.1</td>
<td>ND</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>van't Verlaat, 1991 (44)</td>
<td>Macro</td>
<td>CAB 10.8 mg</td>
<td>12</td>
<td>42.2</td>
<td>33.3</td>
<td>58.6</td>
<td>None</td>
<td>220.0</td>
<td>Tumor volume reduction &gt;50% in all patients</td>
<td>1/12 (8.3%)</td>
<td>12</td>
</tr>
<tr>
<td>Faglia, 1987 (34)</td>
<td>Micro</td>
<td>DHEC 3.0 mg</td>
<td>17</td>
<td>Range 17–49</td>
<td>100</td>
<td>12</td>
<td>3/22 OP, 10/22 BRC</td>
<td>125</td>
<td>7/22 (32%) total disappearance</td>
<td>0</td>
<td>NA</td>
</tr>
<tr>
<td>Liuza, 1985 (37)</td>
<td>Micro</td>
<td>BRC (0.6), 8.4 mg; LIS (4), 0.4 mg</td>
<td>30</td>
<td>42.3</td>
<td>46.7</td>
<td>58.4</td>
<td>5/30 OP + RT, 7/30 OP</td>
<td>1869</td>
<td>In 25/30 (83%) reduction of tumor volume</td>
<td>1/30 (3.3%)</td>
<td>21</td>
</tr>
<tr>
<td>Ho, 1985 (36)</td>
<td>Micro</td>
<td>BRC ?</td>
<td>7</td>
<td>27.4</td>
<td>100</td>
<td>63</td>
<td>2/84/44 OP</td>
<td>85.4</td>
<td>total shrinkage in 4/44</td>
<td>4/7 (57.1%)</td>
<td>28.8</td>
</tr>
<tr>
<td>Winkelmann, 1985 (45)</td>
<td>Micro 5, macro 35</td>
<td>BRC ?</td>
<td>40</td>
<td>ND</td>
<td>52.5</td>
<td>62.4</td>
<td>2/84/44 OP</td>
<td>ND</td>
<td>Total 7/40 (17.5%), micro 45 (80%)</td>
<td>21.5</td>
<td></td>
</tr>
<tr>
<td>Matto, 1985 (40)</td>
<td>Micro</td>
<td>BRC 8.0 mg</td>
<td>32</td>
<td>29.9</td>
<td>100</td>
<td>12</td>
<td>8/36 OP</td>
<td>106</td>
<td>ND</td>
<td>4/2 (12.5%)</td>
<td>24.8</td>
</tr>
<tr>
<td>Mattei, 1984 (38)</td>
<td>Idiopathic</td>
<td>BRC range 5–20 mg^d^</td>
<td>14</td>
<td>ND</td>
<td>100</td>
<td>10.6</td>
<td>None</td>
<td>80.6</td>
<td>NA</td>
<td>3/14 (21.4%)</td>
<td>7.0</td>
</tr>
<tr>
<td>Idiopathic MET range 12–24 mg^d^</td>
<td>8</td>
<td>ND</td>
<td>100</td>
<td>12.9</td>
<td>None</td>
<td>None</td>
<td>57.1</td>
<td>NA</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Idiopathic MET range 12–24 mg^d^</td>
<td>10</td>
<td>ND</td>
<td>100</td>
<td>11.5</td>
<td>None</td>
<td>None</td>
<td>88.9</td>
<td>No reduction in tumor volume</td>
<td>2/10 (20%)</td>
<td>8.0</td>
<td></td>
</tr>
<tr>
<td>Macro</td>
<td>BRC range 5–20 mg^d^</td>
<td>3</td>
<td>ND</td>
<td>18.0</td>
<td>None</td>
<td>None</td>
<td>77.3</td>
<td>No reduction in tumor volume</td>
<td>0</td>
<td>NA</td>
<td></td>
</tr>
</tbody>
</table>

(Continued)
TABLE 1. Continued

<table>
<thead>
<tr>
<th>First author, year of publication (Ref.)</th>
<th>Cause of hyperprolactinemia</th>
<th>Treatment and mean dose (mg/d) (for CAB, mg/wk)</th>
<th>No. of patients</th>
<th>Mean age (yr)</th>
<th>Female sex (%)</th>
<th>Mean treatment duration (months)</th>
<th>PRL before treatment (μg/liter)</th>
<th>Regression of tumor during treatment on MR or CT</th>
<th>Persisting normoprolactinemia, n (%)</th>
<th>Mean follow-up in persisting normoprolactinemia (months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maxson, 1984 (39) Micro</td>
<td></td>
<td>BRC 6.0 mg</td>
<td>5</td>
<td>33.2</td>
<td>100</td>
<td>11.0</td>
<td>211.0</td>
<td>ND</td>
<td>0</td>
<td>NA</td>
</tr>
<tr>
<td>Zarate, 1983 (46) Micro</td>
<td></td>
<td>BRC range 15–20 mg</td>
<td>4</td>
<td>33.3</td>
<td>100</td>
<td>24</td>
<td>129.8</td>
<td>No regression</td>
<td>2/4 (50%)</td>
<td>24</td>
</tr>
<tr>
<td>Coculescu, 1983 (31) Micro</td>
<td></td>
<td>BRC 7.5 mg</td>
<td>2</td>
<td>36.5</td>
<td>100</td>
<td>8.0</td>
<td>146.9</td>
<td>Regression; 6/10 no change</td>
<td>2/4 (50%)</td>
<td>18.8</td>
</tr>
<tr>
<td>Sobrinho, 1981 (43) Macro</td>
<td></td>
<td>BRC 7.5 mg</td>
<td>2</td>
<td>31.0</td>
<td>100</td>
<td>6</td>
<td>3555</td>
<td>NA</td>
<td>0</td>
<td>NA</td>
</tr>
<tr>
<td>Eversmann, 1979 (33) Micro</td>
<td></td>
<td>BRC 3.5 mg</td>
<td>6</td>
<td>32.6</td>
<td>83.3</td>
<td>7.1</td>
<td>74.8</td>
<td>NA</td>
<td>0</td>
<td>NA</td>
</tr>
</tbody>
</table>

BRC, Bromocriptine; CAB, cabergoline; QUI, quinagolide; DHEC, dihydroergocriptine; US, lisuride; MET, metergoline; PRL, prolactin; OP, operation; RT, radiotherapy; NA, not applicable; ND, no data.

- Follow-up duration of those with persisting normoprolactinemia and those with recurrence of hyperprolactinemia.
- Conversion of μU/liter to g/liter by dividing through 30.
- Patient characteristics based on n = 16; 14 of 16 reached normoprolactinemia.
- Patient characteristics based on n = 23 all normoprolactinemic during treatment, but in five no follow-up because of pregnancy.
- Patient characteristics based on n = 11 all normoprolactinemic during treatment, but in one no follow-up because of pregnancy and one because CAB was not withdrawn.
- Patient characteristics based on n = 26; 25 of 26 reached normoprolactinemia.
- Patient characteristics based on n = 12; 11 of 12 reached normoprolactinemia.
- Patient characteristics based on n = 22; 17 of 22 reached normoprolactinemia.
- Patient characteristics based on n = 44; 40 of 44 reached normoprolactinemia.
- Patient characteristics based on n = 36; 32 of 36 reached normoprolactinemia.
- Range of BRC for idiopathic hyperprolactinemia, micro- and macroprolactinomas.
- Range of MET for idiopathic hyperprolactinemia and microprolactinomas.
- Range of BRC for micro- and macroprolactinomas.
- Macroprolactinomas with interruption of the sella turcica on x-ray or clinical evidence of extrasellar expansion of tumor.
- Data for age of n = 8; six of eight reached normoprolactinemia.
from 0 to 74%. This highest proportion of 74% was observed in a series with idiopathic hyperprolactinemia treated with cabergoline. The pooled proportion of patients with persisting normoprolactinemia after dopamine agonist withdrawal was 21% in a random effects model [95% confidence interval (CI), 14–30%; I² 81%].

**Sensitivity analysis (Table 2)**

Restriction of the analysis to four studies using cabergoline as the only treatment showed a pooled proportion persisting normoprolactinemia of 35% (random effects model, 95% CI, 19–56%). In studies using bromocriptine, the proportion persisting normoprolactinemia was lower (20%; 95% CI, 16–26%). Stratified analysis according to cause of hyperprolactinemia showed higher proportions of treatment success in idiopathic hyperprolactinemia (32%; 95% CI, 5–80%) compared with both microprolactinomas (21%; 95% CI, 10–37%) and macroprolactinomas (16%; 95% CI, 6–36%). Higher proportions of persisting normoprolactinemia were shown in studies with treatment duration longer than 24 months (34%; 95% CI, 19–52%), compared with studies with shorter treatment duration (16%; 95% CI, 11–22%). Exclusion from the analysis of two studies that used radiotherapy in some patients (37, 42) showed persisting normoprolactinemia in 39% (95% CI, 34–44%). Studies in which 50% tumor reduction was achieved in all patients before stopping the dopamine agonist showed persisting normoprolactinemia in 55% (95% CI, 36–73%). Excluding two studies with a considerable proportion of pregnant patients during follow-up (29, 42) showed persisting normoprolactinemia in 20% (95% CI, 12–31%). In studies with a prespecified protocol, the treatment success was 18% (95% CI, 9–31%) using a random effects model (I² 84%).

In a random effects meta-regression adjusting for cause of hyperprolactinemia, longer treatment duration was associated with higher proportion of persisting normoprolactinemia (P = 0.015), whereas the use of cabergoline showed a trend of effect (P = 0.07).

Exclusion of the study of Colao et al. (5) from the analysis decreased both the treatment success rates and the heterogeneity for idiopathic hyperprolactinemia (17%; I² 0%), microprolactinomas (19%; I² 39%), and macroprolactinomas (12%; I² 31%). This decreasing heterogeneity showed that from a statistical point of view, the results from that particular study were outliers in relation to the results of the other studies.

**Discussion**

The present systematic review and meta-analysis was performed to estimate the pooled proportion of patients with persistent normoprolactinemia after withdrawal from dopamine agonists. The study showed that withdrawal was associated with persisting normoprolactinemia in only 21% of all patients. Success rates were higher in patients treated for idiopathic hyperprolactinemia, after treatment with cabergoline, and in patients with treatment duration of more than 2 yr.

Randomized controlled studies comparing different withdrawal strategies after successful treatment of hyperprolactinemia are lacking. In 2006, the Pituitary Society provided guidelines as practical clinical tools for the routine clinical care. These guidelines were mainly based on a landmark study by Colao et al. (4) that demonstrates that dopamine agonist treatment indeed can be successfully withdrawn in a considerable proportion of patients, provided that they fulfilled selected clinical criteria, such as significant tumor reduction on radiological imaging and a prolonged period of normoprolactinemia during treatment. The first study that tested the practical applicability of these 2006 Pituitary Society recommendations (6) was
recently published (47). In that study, the estimated 18-month risk of recurrence was 63% in a cohort of 46 selected normoprolactinemic patients previously treated with cabergoline for at least 2 yr. These data exemplify the clinical dilemma: apparently it is indeed possible in routine clinical practice to successfully withdraw selected patients from dopamine agonist treatment, but the likelihood of success is not easily predicted in individual patients.

In agreement, in the present systematic review, the main limitations were the heterogeneity of included patients and treatment regimes. These studies differed markedly with respect to the causes of hyperprolactinemia, the treatment before the start of the dopamine agonists, and the type and duration of dopamine agonist therapy. Despite these sources of heterogeneity, the proportion of patients with persisting normoprolactinemia after withdrawal of dopamine agonists was lower than 30% in the majority of studies. Only three studies reported a higher success rate (5, 36, 46), with a maximum of 74% in patients with idiopathic hyperprolactinemia (5).

It should be noted that the pooled proportion of treatment success is slightly overestimated in the current meta-analysis, because studies with success rates of zero are transformed to avoid zero cells for statistical purposes. Moreover, the pooled proportion is not an accurate reflection of treatment success in all patients with either hyperprolactinemia or prolactinomas, considering that most studies reported withdrawal for only a selected group of patients.

A sensitivity analysis showed that the study with the highest success rate may be viewed as an outlier from a statistical perspective. How does this particular study differ from the remaining ones? Of note, the duration of dopamine agonist treatment was not extremely long compared with other studies. However, the study from Colao et al. (5) differs with respect to two important aspects from all other included studies. First, before withdrawal, the dose was reduced to a minimum level and not abruptly stopped. Second, all patients fulfilling criteria for withdrawal (i.e., tumor regression of >50% on imaging and normoprolactinemia) continued treatment for another 12 months after fulfilling these criteria. In accordance, in another study with persistent normoprolactinemia in more than half of the microprolactinomas, dopamine agonist treatment was explicitly continued several years after normalization of prolactin levels (45). Because that study comprised only seven patients in whom the effect of withdrawal could be adequately assessed, the overall effect of that study for the current meta-analysis was small. A third aspect that could have contributed to the success rates in the study from Colao et al. (5) is the use of cabergoline, which is known to be the most potent of currently available dopamine agonists (48).

One limitation of the study from Colao et al. is that the included patients were clearly selected. From 381 newly diagnosed patients with hyperprolactinemia, 221 (58%) patients were included in the study (5). The patients who were not included can be supposed to have a less favorable outcome with respect to persistent normoprolactinemia after withdrawal of dopamine agonists. This is a limitation for the external validity of the study results (49). The prior probability of a newly diagnosed patient with a prolactinoma that the disease will be in remission after treatment with a dopamine agonist will therefore be lower than the proportions of treatment success reported in that study.

What are the clinical implications of the present meta-analysis? Our study demonstrates that treatment with cabergoline for more than 2 yr is associated with the best outcome. Although it seems reasonable first to reduce the cabergoline dose before withdrawal, this was only protociled in one study (5). In addition, although observational studies have not reported clinical relevant cardiac valve disease after treatment with cabergoline for prolactinomas, the findings obtained with much higher cumulative doses of cabergoline in Parkinson patients underscore that unnecessary prolongation of treatment is undesirable (50–52). Finally, a withdrawal trial in individual patients is unlikely to negatively affect long-term outcome.

In conclusion, this meta-analysis showed that hyperprolactinemia will recur after dopamine agonist withdrawal in a considerable proportion of patients. The probability of treatment success is highest when cabergoline is used for at least 2 yr.

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