

<u>AREA PRESCRIBING COMMITTEE – Birmingham, Sandwell, Solihull and environs</u>

Decision Making Support Tool

The following document supports the committee to consider formulary applications against defined criteria.

| Formulary application reference: | | APCBSSE/0037 | |
|---|--|---------------------|--|
| Drug name and formulations: | | Midodrine (Bramox®) | |
| Criteria | Example | | Committee Consensus |
| Patient Safety | Potential for abuse, toxicity, significant drug interactions | | Number of safety concerns which require regular monitoring and careful management e.g. risk of supine hypertension. |
| Clinical effectiveness | Established licensed product | | Only licensed product for orthostatic hypotension. 2 RCTs found that midodrine significantly increased standing BP 1 hour post-dose compared to placebo. Improvements in patient and investigator-rated symptoms were seen with midodrine compared to placebo. |
| Strength of evidence | | | Moderate, sufficient to get licence. The main limitations of the RCTs was the focus on disease-orientated outcomes (changes in BP), as opposed to patient-orientated outcomes such as quality of life, falls etc. |
| Cost effectiveness or resource impact | £ | | Cost-effective provided the protocol discussed by clinician is followed. |
| Place of therapy relative to available treatments | 1/2 nd tier | | After non-pharmacological intervention: second line pharmacological therapy, in line with licensing. |
| National guidance and priorities | NICE, MTRAC | 2 | NICE Evidence Summary (Oct 2015) |
| Local health priorities | CCG views | | Would support. |
| Equity of access | Equality asse | essment | N/A |
| Stakeholder views | Define wider be engaged | groups to | N/A |
| Implementation requirements | Requires, RIC etc. | CAD ESCA | Requires ESCA but draft put forward with application needs reviewing to include: annual review with specialist, glaucoma monitoring and monitoring interval clarified. |



Decision Summary

| Resubmission is recommended to complete the | |
|---|---|
| information to enable a decision: | |
| Not approved and rationale: | |
| Formulary status (RAG) and rationale | AMBER with ESCA. Rationale: the members felt that due to safety concerns and need for regular monitoring, shared care arrangements were more appropriate than RICaD. |
| Implementation requirements: | |
| Implementation monitoring: | |