## **Midodrine Information Leaflet**

Information for prescribers and pharmacists

#### Indication

• Idiopathic postural hypotension

#### **Doses and Administration**

- The usual starting dose is 2.5mg 2 3 times daily
- The dose should be increased at intervals of 3 5 days until the optimal response is obtained.
- Most patients are controlled on doses at or below 30mg daily in divided doses
- Doses should be taken in the morning, at noon and evening.
- The last dose should be taken at least four hours before bedtime to reduce the risk of supine hypertension
- Some patients may require a morning dose higher than that taken later in the day

## **Pharmacology**

- Midodrine is a directly acting sympathomimetic with selective alphaagonist activity.
- Midodrine enhances smooth muscle tone leading to peripheral vasoconstriction.
- It has no direct cardiac stimulatory effects
- Midodrine slightly decreases cardiac output and renal blood flow. It increases the tone of the internal bladder sphincter and delays the emptying of the bladder.
- Midodrine is a prodrug, the active metabolite is deglymidodrine

### **Contra-indications**

- Midodrine is contraindicated in patients with severe organic heart disease, congestive heart failure, thyrotoxicosis, phaeochromocytoma, acute renal disease, severe renal insufficiency, urinary retention, narrow angle glaucoma, hyperthyroidism, known hypersensitivity to any component of the product
- Patients with a history of CVA should be monitored closely

### Adverse Effects

- Cardiovascular
  - Supine hypertension usually detected at the initiation of therapy and during titration
  - Symptoms of supine hypertension may include: chest pain,
    SOB, palpitations, headache, blurred vision etc
  - o Reflex bradycardia
  - Arrhythmias
- Skin
  - o Rash, pruritis (mainly of the scalp), flushing

- Gastrointestinal
  - o Nausea
  - Vomiting
  - o Dyspepsia
- Central Nervous System
  - o Parathesia
  - o Headache
  - Restlessness
  - Excitability
  - Irritability
- Urinary
  - o Urinary retention

# **Monitoring**

- Supine and standing blood pressures during the use of the drug
- Urea and Electrolytes
- Renal Function
- Heart rate
- Liver Function Tests

# **Drug interactions**

| Cardiac Glycosides                 | Midodrine may potentiate reflex         |
|------------------------------------|---|
|                                    | bradycardia or other kinds of           |
|                                    | conduction disorders and arrhythmia     |
| Alpha and beta adrenergic blocking | The blood pressure raising effects of   |
| drugs                              | midodrine may be antagonised by         |
| _                                  | alpha blockers. The heart rate reducing |
|                                    | effect of beta blockers may be          |
|                                    | increased.                              |
| Atropine                           | Midodrine may enhance the blood         |
|                                    | pressure raising effect of atropine     |
| Tricyclic antidepressants          | Increased sympathomimetic effect        |
| Sympathomimetic agents             | (increase in blood pressure)            |
| Thyroid hormones                   |   |
| MAO-inhibitors                     |   |
| Corticosteroids                    |   |

# **Prescribing Status In Doncaster**

- Red Listed
- Hospital consultants who initiate prescribing will be expected to make arrangements for continuation of therapy and to continue monitoring of the patient

References Summary of Product Characteristics Midon 2010 Martindale: The Complete Drug Reference Patient Information Leaflet Gutron