

## **Annex I**

### **Scientific conclusions and grounds for the variation to the terms of the Marketing Authorisation(s)**

### **Scientific conclusions**

Taking into account the PRAC Assessment Report on the PSUR(s) for mequitazine, the scientific conclusions are as follows:

In view of available data on tremor from spontaneous reports including in some cases a close temporal relationship and positive de-challenge and in view of a known class effect, the PRAC Lead Member State considers a causal relationship between mequitazine and tremor is at least a reasonable possibility. The PRAC concluded that the product information of products containing mequitazine should be amended accordingly.

The CMDh agrees with the scientific conclusions made by the PRAC.

### **Grounds for the variation to the terms of the Marketing Authorisation(s)**

On the basis of the scientific conclusions for mequitazine the CMDh is of the opinion that the benefit-risk balance of the medicinal product(s) containing mequitazine is unchanged subject to the proposed changes to the product information.

The CMDh reaches the position that the marketing authorisation(s) of products in the scope of this single PSUR assessment should be varied. To the extent that additional medicinal products containing mequitazine are currently authorised in the EU or are subject to future authorisation procedures in the EU, the CMDh recommends that the concerned Member States and applicant/marketing authorisation holders take due consideration of this CMDh position.

## **Annex II**

### **Amendments to the product information of the nationally authorised medicinal product(s)**

**Amendments to be included in the relevant sections of the Product Information** (new text **underlined and in bold**, deleted text ~~strike-through~~)

#### **Summary of Product Characteristics**

- Section 4.8

The following adverse reaction(s) should be added under the SOC nervous disorders with a frequency unknown: **tremor**

#### **Package Leaflet**

- Section 4 Possible side effects

The following side effects have also been reported:

#### **Shaking (Tremor)**

### **Annex III**

#### **Timetable for the implementation of this position**

## Timetable for the implementation of this position

Adoption of CMDh position:	October 2020 CMDh meeting
Transmission to National Competent Authorities of the translations of the annexes to the position:	29 November 2020
Implementation of the position by the Member States (submission of the variation by the Marketing Authorisation Holder):	28 January 2021