VPA10983/054/002

Marbocyl P 20 mg Tablets

| Vet - B3 k) Vet - B3 k) Vet - F.II.b.3 a) Vet - B3 k) test (qual in-pr man VRA proc -Ma finis man Man test (qual in-pr man VRA proc -Ma finis man Man Man Man Man Man Man Man M | RA - Vet - B3 k) - k) Deletion of a non-significant in-process (finished product manufacture) - B3 k) Changes to the lity part of the dossier: Deletion of a non-significant rocess test (e.g. deletion of an obsolete test) during the nufacture of the finished product A-R - Vet - F.II.b.3 a) - a) Minor change in the manufacturing cess - F.II.b.3 a) Quality Changes - Finished Product mufacture - Change in the manufacturing process of the shed product, including an intermediate used in the nufacture of the finished product - Minor change in the mufacturing process RA - Vet - B3 k) - k) Deletion of a non-significant in-process (finished product manufacture) - B3 k) Changes to the lity part of the dossier: Deletion of a non-significant | 06/07/23 |
|---|---|----------|
| Vet - F.II.b.3 a) Vet - F.II.b.3 a) Vet - B3 k) Vet - B3 k) | A-R - Vet - F.II.b.3 a) - a) Minor change in the manufacturing cess - F.II.b.3 a) Quality Changes - Finished Product inufacture - Change in the manufacturing process of the shed product, including an intermediate used in the nufacture of the finished product - Minor change in the nufacturing process RA - Vet - B3 k) - k) Deletion of a non-significant in-process (finished product manufacture) - B3 k) Changes to the | 18/05/23 |
| Vet - B3 k) test qual | (finished product manufacture) - B3 k) Changes to the | |
| | rocess test (e.g. deletion of an obsolete test) during the nufacture of the finished product | 11/05/23 |
| Vet - B3 k) test qual in-pu | RA - Vet - B3 k) - k) Deletion of a non-significant in-process (finished product manufacture) - B3 k) Changes to the lity part of the dossier: Deletion of a non-significant rocess test (e.g. deletion of an obsolete test) during the nufacture of the finished product | 09/05/23 |
| Vet - C1 deta doss | RA - Vet - C1 - Change(s) in the name or address or contact iils of a qualified person for pharmacovigilance (QPPV) - C1 nges to the safety, efficacy and pharmacovigilance part of the sier: Change(s) in the name or address or contact details of a lified person for pharmacovigilance (QPPV) | 31/03/23 |
| VNI chan elsev Vet - C6 Chan doss the s | RA - Vet - C6 - Introduction of a summary of the PSMF or nges to the summary of the PSMF not already covered where in the Annex to Regulation (EU) 2021/17 - C6 nges to the safety, efficacy and pharmacovigilance part of the sier: Introduction of a summary of the PSMF or changes to summary of the PSMF not already covered elsewhere in the nex to Regulation (EU) 2021/17 | 31/03/23 |
| A.7 Subs man cont exci ADM sites pack wher reag | A.7 - A.7 Deletion of manufacturing sites for an active stance, intermediate or finished product, packaging site, nufacturer responsible for batch release, site where batch trol takes place, or supplier of a starting material, reagent or pient (when mentioned in the dossier)* - A.7 - MINISTRATIVE CHANGES - Deletion of manufacturing s for an active substance, intermediate or finished product, kaging site, manufacturer responsible for batch release, site ere batch control takes place, or supplier of a starting material, gent or excipient (when mentioned in the dossier)* B.I.d.1.a.4 - 4. Extension or introduction of a re-test | 06/04/22 |

| | period/storage period supported by real time data - B.I.d.1.a.4 - QUALITY CHANGES - ACTIVE SUBSTANCE - Stability - Change in the re-test period/storage period or storage conditions of the active substance where no Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier - Re-test period/storage period - | |
|-----------|--|----------|
| B.I.b.2.e | IB - B.I.b.2.e - e) Other changes to a test procedure (including replacement or addition) for the active substance or a starting material/intermediate - B.I.b.2.e - QUALITY CHANGES - ACTIVE SUBSTANCE - Control of active substance - Change in test procedure for active substance or starting material/reagent/intermediate used in the manufacturing process of the active substance - Other changes to a test procedure (including replacement or addition) for the active substance or a starting material/intermediate | 06/04/22 |
| B.I.b.2.e | IB - B.I.b.2.e - e) Other changes to a test procedure (including replacement or addition) for the active substance or a starting material/intermediate - B.I.b.2.e - QUALITY CHANGES - ACTIVE SUBSTANCE - Control of active substance - Change in test procedure for active substance or starting material/reagent/intermediate used in the manufacturing process of the active substance - Other changes to a test procedure (including replacement or addition) for the active substance or a starting material/intermediate | 06/04/22 |