**VPA22622/023/009**

**Vetoryl 120 mg chewable tablets for dogs**

| Variation | Summary | Date |
|---|---|---|
| Vet - B11 d) | VNRA - Vet - B11 d) - - Vet - B11 d) - Change in the specification parameters or limits of an active substance, starting material, intermediate or reagent used in the manufacturing process of the active substance or of the immediate packaging of the active substance: addition of a new specification parameter to the specification with its corresponding test method for an active substance, starting material, intermediate or reagent used in the manufacturing process of the active substance | 25/08/25 |
| Vet - B11 d) | VNRA - Vet - B11 d) - - Vet - B11 d) - Change in the specification parameters or limits of an active substance, starting material, intermediate or reagent used in the manufacturing process of the active substance or of the immediate packaging of the active substance: addition of a new specification parameter to the specification with its corresponding test method for an active substance, starting material, intermediate or reagent used in the manufacturing process of the active substance | 25/08/25 |
| Vet - B11 d) | VNRA - Vet - B11 d) - - Vet - B11 d) - Change in the specification parameters or limits of an active substance, starting material, intermediate or reagent used in the manufacturing process of the active substance or of the immediate packaging of the active substance: addition of a new specification parameter to the specification with its corresponding test method for an active substance, starting material, intermediate or reagent used in the manufacturing process of the active substance | 25/08/25 |
| Vet - B3 e) | VNRA - Vet - B3 e) - - Vet - B3 e) - Changes to the quality part of the dossier: Deletion of a test procedure — for the active substance or a starting material, reagent or intermediate of the active substance;— for the immediate packaging of the active substance;— for an excipient or the finished product;— for the immediate packaging of the finished product | 25/08/25 |
| Vet - B43 | VNRA - Vet - B43 - - Vet - B43 - Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible | 25/08/25 |
| Vet - B3 d) | VNRA - Vet - B3 d) - - Vet - B3 d) - Changes to the quality part of the dossier: Deletion of a non-significant specification parameter (e.g. deletion of an obsolete parameter) of — an active substance;— a starting material;— an intermediate or reagent used in the manufacturing process of the active substance | 25/08/25 |
| Vet - B3 n) | VNRA - Vet - B3 n) - - Vet - B3 n) - Changes to the quality part of the dossier: Deletion of a non-significant specification | 25/08/25 |

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| | parameter (e.g. deletion of an obsolete parameter such as odour and taste or identification test for a colouring or flavouring material) in the specification parameters or limits of the finished product | |
| Vet - B3 n) | VNRA - Vet - B3 n) - - Vet - B3 n) - Changes to the quality part of the dossier: Deletion of a non-significant specification parameter (e.g. deletion of an obsolete parameter such as odour and taste or identification test for a colouring or flavouring material) in the specification parameters or limits of the finished product | 25/08/25 |
| Vet - F.II.b.5 b) | VRA-S - Vet - F.II.b.5 b) - - Vet - F.II.b.5 b) - Change to in-process tests or limits applied during the manufacture of the finished product - Widening of the approved IPC limits, which may have a significant effect on overall quality of the finished product | 22/08/25 |
| Vet - F.III.1 a) z. | VRA-R - Vet - F.III.1 a) z. - - Vet - F.III.1 a) z. - Submission of a new or updated Ph. Eur. certificate of suitability or deletion of Ph. Eur. certificate of suitability - For an active substance, For a starting material/reagent/intermediate used in the manufacturing process of the active substance, For an excipient - European Pharmacopoeial Certificate of Suitability to the relevant Ph. Eur. Monograph. - Other changes under this code level, e.g. variations outlined in section 6 and 7 of this guidance | 22/08/25 |
| Vet - G.I.15 z) | VRA-R - Vet - G.I.15 z) - z) Other changes under this code level e.g. variations outlined in section 6 and 7 of EMA/CMDv/7381/2021 - G.I.15 z) Safety, Efficacy, Pharmacovigilance changes - Changes to the labelling or the package leaflet which are not connected with the summary of product characteristics - Other changes under this code level, e.g. variations outlined in section 6 and 7 of EMA/CMDv/7381/2021 | 19/11/24 |
| Vet - C10 a) | VNRA - Vet - C10 a) - a) Administrative information concerning the holder’s representative - C10 a) Changes to the safety, efficacy and pharmacovigilance part of the dossier: Changes to the labelling or the package leaflet which shall not be connected with the SPC: — administrative information concerning the holder’s representative | 25/10/24 |