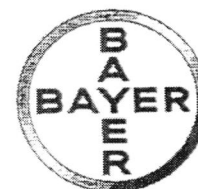


| KVP Pharma+Veterinär Produkte GmbH Kiel, Germany | | Certificate of Analysis | | Page: 1 of 2 Date: 2013-11-15 |
|--|---------------------------|---|-------------|----------------------------------|
| Material: 82258354 Your Material: | | ADVOCATE S.O. FOR SMALL CATS <4KG3X0,4ML BOX X 3 PIP X 0,4 CC ADVOCATE S.O. FOR SMALL CATS < 4 KG | | |
| Batch: | KP0945H | Country: Hungary | | |
| Date of manufacture: | 2013-08-05 | Delivery number: 101391366 | | |
| Expiry date: | 2016-08-31 | Order number: 1880620 | | |
| From Material: | 83196816 | IMI 10%/MOXI 1% K 0,4 ML 80142277 | | |
| Batch: | KP090XZ | Insp. instruction: | T.02.02 - 7 | |
| Inspection lot: | 040001185989 | Specification: | T.02.01 - 4 | |
| Inspection | Acceptance criterion | UoM. | Result | |
| Material (visual) | Solution | | solution | |
| Clarity (visual) | clear | | clear | |
| Colour (Ph.Eur.) | yellow to brownish-yellow | | yellow | |
| Identity (HPLC) | must comply | | complies | |
| Identity (TLC) | must comply | | complies | |
| Assay Imidacloprid | 9.5 - 10.5 | g/hml | 10.0 | |
| Any unspecified degrad. prod.Imidacloprid | max. 1.0 | % | 0.3 | |
| Sum of all degrad. products imidacloprid | max. 1.0 | % | 0.3 | |
| Assay Moxidectin | 0.90 - 1.05 | g/hml | 0.99 | |
| 23-Keto-F-Alpha compound | max. 2.0 | % | < 0.1 | |
| Any unspecified degrad. prod.Moxidectin | max. 1.0 | % | 0.4 | |
| Sum of all degrad. products moxidectin | max. 4.0 | % | 0.4 | |
| Relative density (USP) | 1.093 - 1.103 | | 1.097 | |
| Butylhydroxytoluene | 0.050 - 0.116 | g/hml | 0.103 | |
| Water | max. 6.0 | % | < 0.3 | |



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| Inspection lot: | 040001185989 | | | |
| Inspection | Acceptance criterion | UoM. | Result | |
| Uniformity of content | must comply | | complies | |
| Total aerobic microbial count (TAMC) | max. 100 | CFU/g | *) | |
| Total combined yeast/mould count (TYMC) | max. 10 | CFU/g | *) | |
| Microb.purity Staphylococcus aureus/1g | absent | | *) | |
| Microbial purity Pseudomonas aerug./1g | absent | | *) | |

*) Test is carried out on spot-check basis. However we confirm compliance with the inspection and requirements also for this batch.

This batch complies with the specification and is manufactured in accordance with EC Guide to GMP for Medicinal Products and the requirement laid down in the product license.

Release Documentation is signed by Responsible Person Quality Management - Dr. Amann - QP

This Certificate of Analysis was automatically printed.