

## **Post Authorisation Assessments**

## Octacillin 697 mg/g Powder for Use in Drinking Water for Pigs Vm 16849/4030

r 2021 Update of the test procedure to comply with the updated general Ph. Eur monograph.
Deletion of manufacturing site for an active substance.
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Approval of mock ups.
Update to the SPC and PIL texts according to the latest QRD template.
Submission of a new Ph. Eur. certificate of suitability for
an active substance from a new manufacturer.
Repeat Use application to add 11 new member states.
19 Changes to the DDPS.
Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
r 2018 Submission of a new Ph. Eur. certificate of suitability for
an active substance from a new manufacturer.
Introduction of a re-test period of the active substance.
18 Submission of an updated Ph. Eur. certificate of
suitability for an active substance from an already
approved manufacturer.
18 Submission of an updated Ph. Eur. certificate of
suitability for an active substance from an already
approved manufacturer.
Deletion of manufacturing site for an active substance
manufacturer.
Changes to an existing pharmacovigilance system as
described in the DDPS.
8 Minor change in the manufacturing process of the
finished product.
Submission of an updated Ph.Eur. certificate of suitability
for an active substance from an already approved
manufacturer.
Submission of a new Ph. Eur. certificate of suitability for
an active substance from a new manufacturer.
r 2016 Mock-ups approved.
Change in distributor details
13 Submission of a new Ph. Eur. Certificate of Suitability
Change of name and contact details for QPPV

•	12 October 2012	Changes in the immediate packaging of the finished product.
•	13 June 2012	Submission of an updated certificate of suitability for an already approved manufacturer.
•	26 January 2012	Renewal – UK as CMS