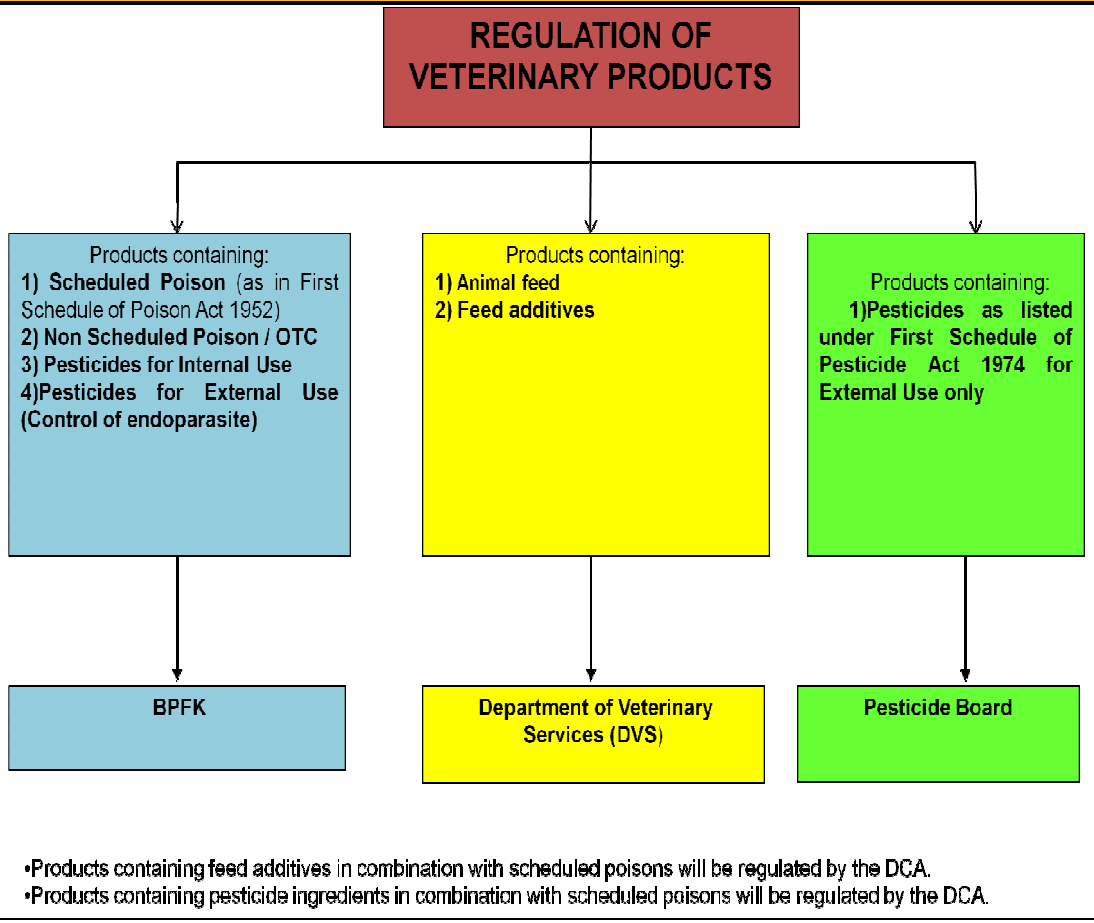
LIST OF UPDATES ON REGOVP, VERSION 3, JULY 2014

<table>
<tr><th rowspan="2">NO.</th><th rowspan="2">REVISION</th><th colspan="2">UPDATES</th><th rowspan="2">REFERENCE</th></tr>
<tr><th>SECTION/ APPENDIX</th><th>DETAILS</th></tr>
<tr><td>1.</td><td>February 2015</td><td>Section E, Inspection, Licensing and Relevant Documents</td><td><u>Amendment at Section E: Inspection, Licensing and Relevant Documents Subsection 13.1: Inspection</u>
<table>
<tr><th>Guidelines</th><th>Product Type/ Category</th></tr>
<tr><td>PIC/S Guide to Good Manufacturing Practice for Medicinal Products *</td><td>Pharmaceuticals (Poison and Non-Poison) Veterinary Products</td></tr>
<tr><td>Guideline on Good Manufacturing Practice (GMP) for Veterinary Premixes; 1st Edition, January 2015</td><td>Veterinary Premixes</td></tr>
<tr><td>Guidelines on Good Distribution Practice (GDP); 2nd Edition 2013</td><td>For activities related to the storage and distribution by manufacturers, importers and wholesalers (where applicable)</td></tr>
</table></td><td>Memo from PKP. Ref: (37)dlm.BPFK/30/06/1 Bhgn 7</td></tr>
<tr><td>2.</td><td>April 2015</td><td>Section A: General Overview</td><td><u>Deletion of Section A: General Overview, Subsection 2.2: (vi)</u></td><td></td></tr>
</table>

NO.	REVISION	UPDATES		REFERENCE
		SECTION/ APPENDIX	DETAILS	
3.	April 2015	Section A: General Overview	**Amendment of Section A: General Overview, Subsection 2.2: (vii) and (viii)**	
4.	April 2015	Appendix 10: Regulation of Veterinary Products in Malaysia	**Amendment of Appendix 10: Regulation of Veterinary Products in Malaysia**	

NO.	REVISION	UPDATES		REFERENCE
		SECTION/ APPENDIX	DETAILS	



•Products containing feed additives in combination with scheduled poisons will be regulated by the DCA.
•Products containing pesticide ingredients in combination with scheduled poisons will be regulated by the DCA.

NO.	REVISION	UPDATES		REFERENCE
		SECTION/ APPENDIX	DETAILS	
5.	July 2015	Section A: General Overview	**Addition of Section A: General Overview, Subsection 2.4**	
6.	July 2015	Section A: General Overview	**Amendment of Section A: General Overview, Subsection 2.6**	
7.	July 2015	Section A: General Overview	**Addition of Section A: General Overview, Subsection 2.2: (x) and (xi)**	

NO.	REVISION	UPDATES		REFERENCE
		SECTION/ APPENDIX	DETAILS	
8.	October 2015	Section A: General Overview Appendix 1: Fees	**Amendment of Section A: General Overview, Subsection 2.5** **Amendment of Appendix 1: Fees, Subsection 1.2**	
9.	August 2016	Appendix 1: Fees Appendix 1.1 – 11 Step 2: New Registration Application Form	**Amendment of Appendix 1: Fees, Subsection 1.4** **Amendment of Numbering of Appendices** **Addition of Section D: Label (Mockup) For Immediate Container, Outer Carton And Proposed Package Insert, Specific Labelling Requirements** **Amendment of Section 2: Guide On How To Fill The Online Application Form For A Product Registration: Check List Of Product Registration Form Entry**	Notice Ref: (40)dlm.BPFK/PPP/01 /03/Jld 3

NO.	REVISION	UPDATES		REFERENCE
		SECTION/ APPENDIX	DETAILS	
10.	November 2016	Appendix 7: List of Ingredients (Active) Not Allowed to Be Registered by The Drug Control Authority	**Amendment of Appendix 7: List of Ingredients (Active) Not Allowed to Be Registered by The Drug Control Authority**	
11.	December 2016	Appendix 7: List of Ingredients (Active) Not Allowed to Be Registered by The Drug Control Authority	**Amendment of Appendix 7: List of Ingredients (Active) Not Allowed to Be Registered by The Drug Control Authority**	