**DEPARTMENT OF PHARMACEUTICAL SERVICES**

**For Official Use:**

**Log Ref. No.: (\_\_\_\_\_)/DRU/Log.Vartn/20\_\_**

**MINISTRY OF HEALTH**

**BRUNEI DARUSSALAM**

**LOG FOR THE APPLICATION FOR VARIATION TO REGISTERED MEDICINAL PRODUCTS**

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| **Name of Product Licence Holder :** | **Date:** |

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| **No.** | **Product Name and Strength** | **Product Licence No.** | **LOA-P/\_/\_\_** | **Variation Code\*** | **FOR OFFICIAL USE** | |
| **Ref.(\_)/DRU/**  **DRA.Variation/20\_** | **Remarks/Query** |
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**\***Please indicate the variation code applied [refer to Appendix 5 – Types of variation of the ‘Guide to Application for Registration of Medicinal Products (4th Edition)’ e.g. MaV-1, MiV-PA1 etc.] If the variation is not listed, please provide the type of variation.

Date Received (For Official Use):