Request for an Expedited Import / Export Licence or Letter of No Objection

Please provide responses to the following questions and return this form to [controlleddrugs@hpra.ie](mailto:controlleddrugs@hpra.ie)

|  |  |
| --- | --- |
| 1. Why is the import or export licence or letter of no objection (LONO) required urgently? |  |
| 1. Please explain why the application was not submitted prior to this, given that the estimated processing time is 4 weeks as outlined in the HPRA ‘Guide to import and export licences for controlled drugs’. |  |
| 1. Will a 4 week processing timeline result in an adverse impact on public health?\* | Yes  No |
| 1. If response to 3 above is yes, please explain why. |  |
| 1. If this application can be expedited, what timeline is desirable to avoid an impact on public health? |  |

**\*** Please note that shortages of an individual brand of a product on a given market may not always have adverse public health implications if alternative brands are available. This should be explored with the appropriate body for that market e.g. Medicines Competent Authority, Health Agency, Department of Health etc. and confirmation provided.

Signed: \_\_\_     \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ Date: \_\_     \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Title: \_\_\_     \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

(Responsible Person/Qualified Person)