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| $\overline{}$ | | | $-\Lambda$ | |

| For official use only | |
|-----------------------|--|
| Application number: | |

Application for "Letter of Notification" of non-medical face mask

| Details of the applicant | | | | |
|--------------------------------|-----------------------------|----------|---------|--|
| 1.Name of the applicant | | | | |
| 2.Address of the applicant | Office address | | | |
| 3.Contact numbers | Residential | | | |
| | Mobile | | | |
| | E mail | | | |
| 4. NIC Number | | | | |
| 5. Type of business | For Local market only | | | |
| | For Export market only | | | |
| | For local and export market | | | |
| Details of the product | | | | |
| 6.Name of the product | | | | |
| 7. Brand name (if any) | | | | |
| 8. Type of material | 100% Cotton 100% Linen | Non- | - woven | |
| Details of the manufacturing s | ite | | | |
| 09. Site address | | Province | | |
| | | District | | |
| | | | | |

Prepared by: Medical Device Regulatory Division & Manufacturing Regulatory Division,

National Medicines Regulatory Authority, Sri Lanka

| 10. Machinery use for | Type of machine/s | | | | | |
|---|--------------------|------|--|--|--|--|
| production | Number if machines | | | | | |
| Make sure following documents are annexed with the application [tick (V)appropriately] | | | | | | |
| Annex I – Finished product test | Yes No | | | | | |
| Annex II – Material confirmation | Yes No | | | | | |
| Annex III – Product label | Yes No | | | | | |
| Annex IV- Colour photograph o | Yes No | | | | | |
| Annex V- Colour photograph o | Yes No | | | | | |
| Annex VI – Copy of Business Re | Yes No | | | | | |
| I,, the undersigned, hereby declare that all the information submitted with this application is true and correct and certify that all documents annexed in support of this application are accurate and correct. | | | | | | |
| I agree to provide any further information and documents requested by the National Medicines Regulatory Authority, which is required for processing of this application and, facilitate any requests by the National Medicines Regulatory Authority for inspection of premises and/or records. I also agree to provide samples of product(s) required for evaluation and/or analysis. | | | | | | |
| I further declare that I take full responsibility for all consequences, which might arise from false or erroneous information submitted in the application and that I will cooperate with any official of the National Medicines Regulatory Authority for any such investigations relevant to the application. | | | | | | |
| | | | | | | |
| Signature of the applicant | | Date | | | | |

Prepared by: Medical Device Regulatory Division & Manufacturing Regulatory Division, National Medicines Regulatory Authority, Sri Lanka

Version 1.0 08th June 2020