AGENDA FOR THE 116th UAC MEETING (VSEZ UNITS) SCHEDULED AT 1100 A.M ON 05.05.2020 UNDER THE CHAIRMANSHIP OF DEVELOPMENT COMMISSIONER, VSEZ, SHRI ARAMA MOHAN REDDY, IFS, IN THE CONFERENCE HALL OF ADMINISTRATIVE BUILDING, DUVVADA

Agenda item No.1.

Ratification of the minutes of the UAC meeting for VSEZ units held on 13.03.2020.

Agenda item No.2

Ratification of issuance of Letter of Approval (LoA) to M/s. Khanija Recycling India Pvt.Ltd., Bangalore – 562107 for setting up of a unit under trading activity for Shredded Printed Circuit Boards (under ITC HS code 84716090) in VSEZ, Duvvada, Visakhapatnam.

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The issue was discussed in the UAC on 13.3.2020 and the UAC decided the matter to be dealt on file. The matter has been dealt on file and Letter of Approval (LoA) dt.27.05.2020 issued to M/s. Khanija Recycling India Pvt.Ltd., with the standard conditions prescribed by DoC and further following terms and conditions:

(i) They shall be permitted to export the permitted goods under this Letter of Permission to Japan only as the item falls under Free and the Government of Japan do not have any restrictions on import of scrap of Printed Circuit Boards.

(ii) They shall, as per Rule 14(1) of Hazardous and other Wastes (Management and Transboundary Movement) Rules, 2016, make an application in Form 5 along with insurance cover to the Ministry of Environment, Forest and Climate Change, Govt. of India for the proposed trans boundary movement of Hazardous and other wastes together with the prior informed consent in writing from the importing country in respect of wastes specified in Part A of Schedule III and Schedule VI of the said Rules.

Also it is opined by the Specified Officer, Customs, VSEZ that Part B of Schedule III Electrical Waste and electronic assemblies or scrap including ‘Printed Circuit Boards’ in Serial No. B 1110 is under the list of other wastes, applicable for import and export and not require prior informed consent of the importing country.

(iii) They shall be required to take permission of State Pollution Control Board and also permission from the Ministry of Environment, Forest and Climate Change for the purpose of export, wherever required.

The matter is placed before UAC for ratification of the LoA issued to the Unit.

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Agenda item No:3

Ratification of approval granted to M/s. Reddy’s Laboratories Ltd., Unit VII, VSEZ for inclusion of additional product in the LoA

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The request of the unit for inclusion of the following products in their LoA under broad banding

<table>
<thead>
<tr>
<th>Sl.No.</th>
<th>Item of manufacture</th>
<th>product</th>
<th>ITCHS</th>
<th>Annual quantity Nos.</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Pazopanib HCL Tablets 200 mg</td>
<td>Tablet</td>
<td>30049049</td>
<td>576921</td>
</tr>
<tr>
<td>2</td>
<td>Pazopanib HCL Tablets 400 mg</td>
<td>Tablet</td>
<td>30049049</td>
<td>192919</td>
</tr>
</tbody>
</table>

Further, the unit submitted a copy of test licence in form 29 for manufacture for the purpose of examination, test for chemical and instrumental analysis and are not for any commercial use and shall be used for bio/clinical studies subject to the grant of BE/CT permission from the DCGI office, and the licence will be valid for a period of three years from its issue dt.14.11.2019

In view of the prevailing conditions due to Covid 19 pandemic, Office of Development Commissioner considered their request on file and since the unit submitted Test Licence and declared the products are not prohibited/restricted, approval granted in terms of para 19(2) of SEZ Rules, 2006 for inclusion of new products in the existing LoA subject to ratification by UAC.

In view of the above, the approval granted by DC’s office placed before UAC for ratification.

Agenda item No.4

Ratification of approval granted to M/s. Reddy’s Laboratories Ltd., IX, VSEZ for inclusion of additional product in the LoA

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The unit was issued LoA dt.08.06.2012 for manufacture of pharmaceutical tablets, capsules, injections and commenced production w.e.f.1.11.2017. The unit intends to add the following new products in their LoA:

<table>
<thead>
<tr>
<th>Sl.No.</th>
<th>Item of manufacture</th>
<th>Injections/Vial</th>
<th>ITCHS</th>
<th>Exhibit quantity in Nos.</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Sterile Diluent for Treprostinil Inj</td>
<td>Injection/vial</td>
<td>30041010</td>
<td>6000</td>
</tr>
</tbody>
</table>

Contd.. p 3
The unit submitted photo-copy of Test Licence 623/P&B/2020 dt.13.02.2020 from Drug Control Administration, Govt.of A.P. for the purpose of Examination, Test for chemical and instrumental analysis and not for any commercial use and shall be used for Bio/clinical studies subject to grant of BE/CT permission from the DGG(I) office.

In view of the prevailing conditions due to Covid 19 pandemic, Office of Development Commissioner considered their request on file and since the unit submitted Test Licence and declared the products are not prohibited/restricted, approval granted in terms of para 19(2) of SEZ Rules, 2006 for inclusion of new products in the existing LoA subject to ratification by UAC.

In view of the above, the approval granted by DC’s office placed before UAC for ratification.

Agenda item No.5 /\

Ratification of approval granted to M/s. Shintex Apparel Pvt. Ltd., by office of DC, VSEZ for broad banding and amendment of product approved under broad banding

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The unit is holding LoA dt.08.08.2016 for Laminated Fabric/fabrics for seat cover and Met-T shirt and Trading of laminated fabric/fabrics /\

The unit was issued approval and permission to manufacture and export of ‘PPE Kit’ for trading activity vide letter dt.19.05.2020 under broad banding and also for Body suit, face shield, shoe cover, FFP mask with and without valve for manufacture and export and for trading activity Hand gloves, Sanitizers, N-95 masks, FFP Mask with end without valve vide letter dt.14.04.2020.

Further amendment issued, vide letter dt.19.05.2020, to the product, ‘Body suit’ as PPE KIT that was issued under broad banding vide letter dt.14.04.2020 on file, due to non-convening of meetings of UAC in view of Covid 19 pandemic.

The permission granted to the unit is placed before UAC for ratification.

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Agenda item No.6

Ratification of permission granted to M/s. Lee Pharma Ltd., for manufacture of 'Liquid Hand Sanitizer' under broad banding – reg.

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The unit is holding LoA dt. 30.5.2006 for manufacture of 'Delayed release pellets and granules.

In view of the demand for Sanitizers due to Covid 19 pandemic, the unit was issued permission on file under broad banding for manufacture and export of “Liquid Hand Sanitizer” vide permission letter dt. 07.04.2020 issued via e-mail to the unit.

The permission granted to the unit is placed before UAC for ratification.

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