22 February 2012

GMP COMPLIANCE DECLARATION

To whom it may concern

I, the undersigned, Judith Steinbach, Manager, Bristol-Myers Squibb Quality Operations External Manufacturing, North America declare that the product, Plastibase, manufactured at Contract Pharmaceuticals Limited in Mississauga, Canada, is manufactured under a validated process in compliance with the current Good Manufacturing Practices.

Judith Steinbach
Manager, Quality Operations External Manufacturing, North America
Bristol-Myers Squibb Company
Establishment Licence

CONTRACT PHARMACEUTICALS LTD. CANADA

7600 DANBRO CRESCENT
MISSISSAUGA, ON, CANADA, L5N 6L6

This licence is issued in accordance with the Food and Drugs Act & Regulations (Division 1A & 2) for the following activities and categories of drugs:

<table>
<thead>
<tr>
<th>ACTIVITY/ Catégorie</th>
<th>Pharmaceutical Prod. pharmaceutique</th>
<th>Vaccins</th>
<th>Blood</th>
<th>Schedule D</th>
<th>Schedule C</th>
<th>6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fabricate Manufacturer</td>
<td>X</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Package / label Emballer-étiqueter</td>
<td>X</td>
<td></td>
<td></td>
<td></td>
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<td></td>
</tr>
<tr>
<td>Test Analyser</td>
<td>X</td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Distribute Distribuer</td>
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<td></td>
</tr>
<tr>
<td>Import Importer</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Wholesale Vendre en gros</td>
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<td></td>
<td></td>
</tr>
</tbody>
</table>

(1) Perform the tests, including any examinations required under Division 2 / Analyser conformément au titre 2
(2) Distribute as set out in paragraph C.01A.003 (a) and/or (b) / Distribuer à titre de distributeur au sens de l'alinéa C.01A.003 (a) et/ou (b)
(3) Whole blood and its components / Sang entier et ses composants
(4) Drugs listed in Schedule D to the Act, other than vaccines or whole blood and its components / Drogue visée à l'annexe D de la Loi, autre qu'un vaccin ou le sang entier et ses composants
(5) Drugs listed in Schedule C to the Act / Drogue visée à l'annexe C de la Loi
(6) Drugs listed in the Schedule I Part G of the Food and Drug Regulations, drugs listed in Schedule F to the Food and Drug Regulations, narcotics as defined in section 2 of the Narcotic Control Regulations / Drogue visée à l'annexe Partie G des Règlements sur les aliments et drogues, drogue visée à l'annexe F des Règlements sur les aliments et drogues, stupéfiants au sens de l'article 2 des Règlements sur les stupéfiants
(7) S refers to sterile dosage forms / S fait référence aux formes posologiques stériles

This licence is subject to the additional conditions as indicated in the attached:

Issued On / Emise le: 2010-07-22

Last Inspection Date / Date de la dernière inspection: 2009-08-11

MINISTER OF HEALTH

Stéphanie Reid

Compliance Coordination & Licensing Division

AUG 06 2010

This licence is the property of the Health Products and Food Branch Inspectorate and must be returned upon demand.

This licence appartient à l'Inspectorat de la Direction générale des produits de santé et des aliments et doit être retournée sur demande.
**Establishment Licence**

**CONTRACT PHARMACEUTICALS LTD. CANADA**

1 VALLEYWOOD DRIVE, UNIT 100

MARKHAM, ON, CANADA, L3R 5L9

This licence is issued in accordance with the Food and Drugs Act & Regulations (Division 1A & 2) for the following activities and categories of drugs:

<table>
<thead>
<tr>
<th>ACTIVITY/Activité</th>
<th>CATEGORY/ Catégorie</th>
<th>Pharmaceutical Prod. pharmaceutique</th>
<th>Vaccines</th>
<th>Blood</th>
<th>Schedule D</th>
<th>Schedule C</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fabricate/Manufacturer</td>
<td></td>
<td>(S - Sterile) 7</td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Package/label/Emballer-étiqueter</td>
<td></td>
<td></td>
<td></td>
<td>S</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Test/Analyser</td>
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<td></td>
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</tr>
<tr>
<td>Distribute/Distribuer</td>
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<td></td>
</tr>
<tr>
<td>Import/Importer</td>
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</tr>
<tr>
<td>Wholesale/Vendre en gros</td>
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<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

(1) Perform the tests, including any examinations required under Division 2 / Analyser conformément au titre 2
(2) Distribute as set out in paragraph C.01.A.003 (a) and/or (b) / Distribuer à titre de distribuer au sens de l’alinéa C.01.A.003 (a) et/ou (b)
(3) Whole blood and its components / Sang entier et ses composants
(4) Drugs listed in Schedule D to the Act, other than vaccines or whole blood and its components / Drogue visée à l’annexe D de la Loi, autre qu’un vaccin ou que le sang entier et ses composants
(5) Drugs listed in Schedule C to the Act / Drogue visée à l’annexe C de la Loi
(6) Drugs listed in the Schedule to Part G of the Food and Drug Regulations, drugs listed in Schedule F to the Food and Drug Regulations, narcotics as defined in section 2 of the Narcotic Control Regulations / Drogue visée à l’annexe de Partie G des Règlements sur les aliments et drogues, drogue visée à l’annexe F des Règlements sur les aliments et drogues, stupéfiants au sens de l’article 2 des Règlements sur les stupéfiants
(7) S refers to sterile dosage forms / S fait référence aux formes posologiques stériles

This licence is subject to the additional conditions as indicated in the attached:

Issued On / Emise le: 2010-12-08

Last Inspection Date / Date de la dernière inspection: 2009-12-01

MINISTER OF HEALTH

MINISTRE DE LA SANTÉ

Director Compliance Coordination & Licensing Division

This licence is the property of the Health Products and Food Branch Inspectorate and must be returned upon demand.

Ce licence appartient à l'inspecteur de la Direction générale des produits de santé et des aliments et doit être retournée sur demande.
07 February 2012

To Whom It May Concern:

Subject: Statement on the Transmissible Spongiform Encephalopathies (TSE) status of Plastibase manufactured by Contract Pharmaceuticals Limited (CPL) for Bristol-Myers Squibb Company

The Bristol-Myers Squibb Company (BMS) hereby certifies that Plastibase manufactured at CPL in Mississauga, Ontario, Canada for Bristol-Myers Squibb Company meet the criteria for acceptability based upon an evaluation of TSE risk conducted in accordance with the BMS Corporate Policy and Program for the Assessment of Risk of Transmissible Spongiform Encephalopathies Arising From the Use of Animal-Derived Materials.

BMS evaluated all the raw materials used in the manufacture of Plastibase. All excipients used in the manufacture of Plastibase, manufactured by CPL, were also evaluated.

- All raw materials used in the manufacture of Plastibase are non animal-derived materials

Susan Manne

Susan Manne
Associate Manager, Regulatory & Compliance - IPR/TSE
Research & Development – GRS-CMC
Bristol-Myers Squibb Company
February 7, 2012

Subject: Plastibase®

Dear Customer:

Please note that neither the ingredients nor the manufacturing process of Plastibase® (Polyethylene – Mineral Oil Gel Base) contain or introduce any of the solvents referenced as Class 1, 2, and 3 as described in the current USP 34 chapter <467> Residual Solvents.

According, Plastibase® (Polyethylene – Mineral Oil Gel Base) is not tested for residual solvents and is in alignment with the requirements with the current USP 34 chapter <467> Residual Solvents.
North American Free Trade Agreement

CERTIFICATE OF ORIGIN

1. Exporter's Name and Address:
   Contract Pharmaceuticals Limited
   7600 Danbro Cres.
   Mississauga, ON L5N 6L6

   Tax Identification No: 123057713RMM001

2. Blanket Period:
   From: 01/01/12 To: 12/31/12

3. Producer's Name and Address:
   Contract Pharmaceuticals Limited
   7600 Danbro Cres.
   Mississauga, ON L5N 6L6

   Tax Identification No: 123057713RMM001

4. Importer's Name and Address:
   Bristol Myers Squibb
   777 Scudders Mill Road
   Plainsboro, NJ 08536

5. Description of Good(s):
   Plastibase, Bulk

6. H.S. Tariff
   Classification Number: 2710.19
   Preference Criterion: B
   Producer: YES
   Net Cost: NO
   Country of Origin: Canada

7. I Certify That:
   - The information on this document is true and accurate and I assume the responsibility for proving such representations, I understand that I am liable
   for any false statements or material omissions made on or in connection with this document;
   - I agree to maintain and present upon request documentation necessary to support this certificate, and to inform, in writing, all persons to whom the certificate
   was given of any changes that could affect the accuracy or validity of this certificate;
   - The goods originated in the territory of one or more of the Parties, and comply with the origin requirements specified for those goods in the North
   American Free Trade Agreement, and unless specifically exempted in Article 411 or Annex 401, there has been no further production or any other
   operation outside the territories of the Parties, and
   - This certificate consists of 1 page, including all attachments.

Authorized Signature: [Signature]
Name (Print or Type): Kathleen Smith
Title: Traffic Co-ordinator
Date: 01/01/2012
Telephone: 905-821-7600
Fax: 905-821-7601