様式第十八(第三十六条、第百三十七条の十九関係)

Form No. 18 (related to Article 36 and Article 137—19)

収入印紙 revenue stamp 医 薬 品 医薬部外品外国製造業者 認定申請書 再生医療等製品

drug

Application for accreditation of foreign

quasi-drug regenerative, cellular therapy and gene therapy

manufacturer

products

| 製造所の名称   |          |
|--|----------|
| Name of the manufacturing establishment  |          |
| 製 造 所 の 所 在 地  |          |
| Location of the manufacturing establishment   ②  |          |
| Accreditation categories   |          |
| 製造所の構造設備の概要  |          |
| Outline of the buildings and facilities of   |          |
| the manufacturing establishment  |          |
| 製造所の責任者  |          |
| The person responsible for the Name  |          |
| manufacturing establishment 住所 Address   |          |
|  | いな       |
| C r Ap 条甲 r S P 項請 い者 Applicant whose accreditation was canceled pursuant to the provision of Ar Applicant whose accreditation was canceled pursuant to the provision of Ar e 75—4, Paragraph 1 and who is awaiting a lapse of 3 years from the date said rescission (2) 注第75条の5第1項の規定により認定を取り得され、取消しの日から3年を終過して   |          |
| Applicant whose accreditation was canceled pursuant to the provision of Ar   |          |
| e 75—4, Paragraph 1 and who is awaiting a lapse of 3 years from the date   | of       |
| c r A 条申 rpp 項請 い者 Applicant whose accreditation was canceled pursuant to the provision of Ar attion   | 1,172    |
|  | ۷ ./۲    |
| Applicant whose registration was canceled pursuant to the provision of Art   | icle     |
| To-5, Paragraph 1 and who is awaiting a lapse of 3 years from the date   | of s     |
| I - L and recenterion  | 5// 0    |
| s i. c and rescrission (3) 禁錮以上の刑に処せられ、その執行を終わり、又は執行を受けることがなくなつた行<br>v. c 事 年を経過していない者 Applicant who has a history of a court sentence of imprisonment on sever  | 发、3      |
| Applicant who has a history of a court sentence of imprisonment on seven   | rer      |
| punishment and has not passed 3 years since the execution was completed or   |          |
| longer received  |          |
| (4) 法、麻薬及び向精神薬取締法、毒物及び劇物取締法その他薬事に関する法令で政令<br>の   |          |
| │ arc 3g   |          |
| a d l l l l l l l l l l l l l l l l l l  | ics      |
| tion 任 Control Law, Poisonous and Deleterious Substances Control Law or other 1  |          |
| and regulations related to pharmaceutical affairs specified by Cabinet Or  | der      |
| and regulations related to pharmaceutical affairs specified by Cabinet Or and has not passed 2 years since its date of the disposition  (5) 麻薬 大麻 あへん又は覚醒剤の中毒者   |          |
|  |          |
|  | 判除       |
| r <sub>s +</sub> 員   (6) 精神の機能の障害により外国製造業者の業務を適正に行うに当たつて必要な認知、<br>  <sub>a.e</sub> を   及び意思疎通を適切に行うことができない者  | 1.01-541 |
| Applicant who cannot properly perform the necessary recognition, judgem  |          |
| Applicant who cannot properly perform the necessary recognition, judgem and communication to perform the work of foreign manufacturers properly to mental dysfunction  | due      |
| e ii ∪ to mental dysfunction o i: の (7) 外国製造業者の業務を適切に行うことができる知識及び経験を有すると認められた  | 1 1 本    |
| The transfer of the state of th |          |
| Applicant who is not recognized as naving knowledge and experience to properly carry out the work of foreign manufacturers   |          |
| 備考   |          |
| Remarks  |          |

医 楽 品 上記により、医薬部外品の外国製造業者の認定を申請します。 再生医療等製品

drug

I hereby apply for the accreditation of the foreign quasi-drug

regenerative, cellular therapy and gene

manufacturer indicated above.

therapy products

年 月 日 Year Month Day

邦文

Japanese

住 所 Address

外国文

Foreign language

法人にあつては、主たる事務所の所在地

Location of the head office in case of a corporation

邦文 Japanese

氏 名 外国文

Name Foreign language

法人にあつては、名称及び代表者の氏名

Name and name of its representative in case of a corporation

厚生労働大臣 殿

To Minister of Health, Labour and Welfare

## (注意)

(Notes)

1 用紙の大きさは、A4とすること。

Use paper of Japanese Industrial Standards Size A4.

2 この申請書は、正副2通提出すること。

Applicant should submit one original and one copy of it.

3 字は、墨、インク等を用い、邦文にあつては、楷書ではつきりと書くこと。

Fill in the form with clear writing with inks, etc.,.

4 収入印紙は、正本にのみ貼り、消印をしないこと。

Put revenue stamp only on the original and do not cancel it.

5 認定の区分欄には、第35条第1項及び第2項各号又は第137条の18各号のいずれに該当するかを記載すること。

Identify in the column of "Accreditation categories" which category specified under Article 35, Paragraph 1 and 2 or Article 137—18 is applied.

6 製造所の構造設備の概要欄にその記載事項の全てを記載することができないときは、同欄に「別紙のとおり」と記載し、別紙を添付すること。

In case there is not enough space to fill in all the information in the column "Outline of the buildings and facilities of the manufacturing establishment", write "see attached paper" in the column and attach another paper on which all the information is written.

7 申請者の欠格条項の(1)欄から(7)欄までには、当該事実がないときは「なし」と記載し、あるときは、(1)欄及び(2)欄にあつてはその理由及び年月日を、(3)欄にあつてはその罪、刑、刑の確定年月日及びその執行を終わり、又は執行を受けることがなくなつた場合はその年月日を、(4)欄にあつてはその違反の事実及び違反した年月日を記載すること。

Describe "No" in each column of (1), (2), (3), (4), (5), (6) and (7) if an applicant doesn't meet any conditions of its disqualifications. If an applicant meets one or more conditions of its disqualifications, describe as below.

Column (1) and (2): The date (year, month, day) and its ground for the cancellation.

Column (3) : The date (year, month, day) of final judgment of the crime, sentence and the date (year, month, day) of the completion of its execution.

: The fact and the date (year, month, day) of its violation(s).

Column (4)