|  |
| --- |
| 様式２ |
|  |

医薬品

医薬部外品

再生医療等製品

変更

追加

|  |  |  |  |
| --- | --- | --- | --- |
|  | 区分 |  | 認定書 |
| Certificate of accreditation on category　　　　　for foreign manufacturer  (drug / quasi-drug / regenerative, cellular therapy and gene therapy products)  change  addition | | | |

|  |  |
| --- | --- |
| 氏名又は名称  Name (Name of  corporation) |  |

|  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| 変更  追加 | |  | | 年 |  | 月 |  | 日付けで申請のあった | | | |
| 区分の |  | | を医薬品、医療機器等の品質、有効性及び安全性の確保等に  第13条の3第3項  第23条の24第3項 | | | | | | | | |
| 関する法律（昭和35年法律第145号）　　　　　　　　において準用する　　　　　　　　　　　の規定により、申請のとおり認定する。  Article 13, Paragraph 8  Article 23-22, Paragraph 8  第13条第8項  第23条の22第8項 | | | | | | | | | | | |
| In accordance with the provision of 　　　　　　　　　　　　applied corresponding to  Article 13-3, Paragraph 3  Article 23-24, Paragraph 3 | | | | | | | | | | | |
| of the Act on Securing Quality, Efficacy and Safety of | | | | | | | | | | | |
| Pharmaceuticals, Medical Devices, Regenerative and Cellular Therapy Products,  change  addition | | | | | | | | | | | |
| Gene Therapy Products, and Cosmetics (Act No. 145, 1960), the application for | | | | | | | | | | |  |
| in accreditation category of the foreign manufacturer dated | | | | | | | | |  | is accredited | |
| as applied. | | | | | | | | | | | |

|  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | |  | |  | 年 |  | 月 |  | 日 |
|  | |  | | Year | | Month | | Day |
|  | |  | |  | |  | |  |

|  |  |  |
| --- | --- | --- |
| 厚生労働大臣  Minister of Health, Labour and Welfare |  |  |
|  | | |
|  | | |
|  | | |
|  | | |
|  | | |