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| 様式２　 |
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医薬品

医薬部外品

再生医療等製品

変更

追加

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

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| 氏名又は名称Name (Name ofcorporation) |  |

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| 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 8Article 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 toArticle 13-3, Paragraph 3Article 23-24, Paragraph 3 |
| 　　　　　　　　 　of the Act on Securing Quality, Efficacy and Safety of |
| Pharmaceuticals, Medical Devices, Regenerative and Cellular Therapy Products,changeaddition |
| Gene Therapy Products, and Cosmetics (Act No. 145, 1960), the application for |  |
| in accreditation category of the foreign manufacturer dated |  | is accredited |
| as applied. |

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