Medicines & Healthcare products Regulatory Agency

MHRA

Our Ref: IVD001159

Mr Peter Wei
Lotus Global co Ltd
1 Fourseasons Terrace
West Drayton
Middlesex
UB7 9GG
United Kingdom

MHRA
10 South Colonnade
Canary Wharf
London
E14 4PU
United Kingdom

05 May 2020

Dear Mr Wei

IN VITRO DIAGNOSTIC MEDICAL DEVICES REGULATIONS 2002: REGULATION 44

Registration of manufacturers of *In-Vitro Diagnostic* Medical Devices and devices for Performance Evaluation

Thank you for informing the Competent Authority of the details of ***Manufacturers*** **Name:- MULTIBRANDS TRADING (SUZHOU) CO., LTD** located at **Manufacturers Address:- Room 313, Ruiyuan Building No. 58, Dongda Street, Suzhou China 215002** for whom you are acting as the authorised representative and for supplying the medical device information.

Your registration has been recorded based on your declaration that you have determined that the device(s) fall within the definition of "in vitro diagnostic medical device", and that you have classified it/them correctly considering the intended purpose(s) and mode(s) of action. In accepting your registration, I should make clear that the Competent Authority does not examine each individual notification and therefore cannot and does not necessarily endorse these determinations.

Please note this letter does not represent any form of accreditation, certifcation or approval by the UK Competent Authority.

If you stop placing devices on the market or if you are not complying with the Regulations you should inform us as required by the Regulations. You should be aware that it is an offence to place on the market CE marked devices that do not comply with the Regulations.

The information you provided has been recorded against the reference number shown at the top of this letter, which we ask you to quote in all future correspondence and communications.

Please inform us of any of the following changes;

- the company information
- additional generic groups of devices or, for Annex II or Self-Test devices, additional devices
- discontinuation of a generic group of devices or, for Annex II or Self-Test devices, discontinuation of devices

You should submit your change of registration via DORS with the required statutory fee, which should be accompanied with the information when it is supplied, (the fee is payable for each record notified, and you may place multiple changes on one record).

Thank you for registering the following generic groups of devices

1. ***Part 5: IVDs which are not Annex II and not self-test devices***
2.

3. For reagnets, reagent products, calibration and control materials:
4. group by common technological characteristics and/or analytes
5.
6. New products:
7. Coronavirus
8. Coronavirus - NA Reagents
9.
10. For performance evaluation:
11. None
12.
13. Neither:
14. None
15.
16.
17. For other IVDs, group by appropriate indications
18.
19. New products:
20. None
21.
22. For performance evaluation:
23. None
24.
25. Neither:
26. None
27.
28.
29. Part 6: IVDs which are Annex II or self-test devices
30.
31. For reagnets, reagent products, calibration and control materials:
32. group by common technological characteristics and/or analytes
33.
34. New products:
35. None
36.
37. For performance evaluation:
38. None
39.
40. Neither:
41. None
42.
43.
44. For other IVDs, group by appropriate indications
45.
46. New products:
47. None
48.
49. For performance evaluation:
50. None
51.
52. Neither:
53. None
54.

If you have any queries regarding your registration, please do not hesitate to contact us.

Yours sincerely

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Malcolm Ridgway
Data Integrity Support Officer