

Certificate of Registration



This is to certify that the Occupational Health & Safety Management System of:

Track Components Ltd

Unit 3a, Corinium Industrial Estate, Raans Road, Amersham, Buckinghamshire, HP6 6JQ, United Kingdom

(Central function listed above. See appendix for additional locations)

applicable to:

The Health & Safety management of the design ,development, manufacture of precision CNC turned and milled components, and the provision of sub assembly and assembly capability

has been assessed and registered by NQA against the provisions of:

ISO 45001:2018

This registration is subject to the company maintaining an occupational health & safety management system, to the above standard, which will be monitored by NQA.



A handwritten signature in black ink, appearing to read 'M Winger', is positioned above the title 'Managing Director'.

Managing Director

Certificate No.	187552
ISO Approval Date:	12 June 2023
Valid Until:	11 June 2026
EAC Code:	17, 19, 29



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Certificate of Registration



Appendix to Certificate Number: 187552

Includes Facilities Located at:

Track Components Ltd

Certificate No. 187552
Unit 3a, Corinium Industrial Estate
Raans Road
Amersham Buckinghamshire HP6 6JQ
United Kingdom

The health and safety management of all activities and processes for the manufacture of precision CNC turned and milled components, and the provision of sub assembly and assembly capability

Arcadia Desiccant Technology

Certificate No. 187552/1
Unit 3A Corinium Industrial Estate Raans Road
Amersham Buckinghamshire HP6 6JQ
United Kingdom

The health and safety management of all activities and processes for the design, development, manufacture and assembly of environmental control equipment, including the sale and distribution of associated product

Track Medical

Certificate No. 187552/2
Unit 3a, Corinium Industrial Estate
Raans Road
Amersham Buckinghamshire HP6 6JQ
United Kingdom

The health and safety management of all activities and processes for the provision of assembly services to manufacturers within the medical devices industry, in accordance with their specifications, and applicable regulatory requirements



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