

Commission Regulation (EU) No 383/2012 of 4 May 2012 laying down technical requirements with regard to driving licences which include a storage medium (microchip) (Text with EEA relevance)

Article 1	Scope
Article 2	General requirements
Article 3	Applicable standards
Article 4	Procedure for EU type-approval
Article 5	EU type-approval certificate
Article 6	Single contact points
Article 7	Safeguard clause
Article 8	Entry into force
	Signature

ANNEX I

General requirements for driving licences which include a microchip

The general requirements for driving licences including a microchip described...
the specifications for the microchip and the logical data structure...

- I.1 ABBREVIATIONS
- I.2 DATA STORED ON THE MICROCHIP
 - I.2.1 Harmonised mandatory and optional driving licence data
 - I.2.2 Additional data
- I.3 MICROCHIP
 - I.3.1 Storage medium type
 - I.3.2 Applications
 - I.3.2.1 EU Driving Licence Application
 - I.3.2.2 Other Applications
- I.4 LOGICAL DATA STRUCTURE OF THE EU DRIVING LICENCE APPLICATION
 - I.4.1 Logical Data Structure
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- I.5 DATA PROTECTION MECHANISMS
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 - I.5.1.2 Optional Active Authentication
 - I.5.2 Access Restriction
 - I.5.2.1 Mandatory Basic Access Protection
 - I.5.2.2 Conditional Extended Access Control

I.5.3 Public Key Infrastructure (PKI) for driving licences including a microchip...

- I.6 DATA PRESENTATION
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ANNEX II

ANNEX III

Procedure for EU type-approval of driving licences which include a microchip

- III.1 GENERAL PROVISIONS
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III.3 FUNCTIONAL CERTIFICATE
III.4 EU TYPE-APPROVAL CERTIFICATE
III.4.1 Model certificate
III.4.2 Numbering system

ANNEX IV

Model for the EU type-approval certificate concerning driving licences which include a microchip

Name of the competent authority: ...
Notification concerning:
approval # withdrawal of approval #
of an EU Driving Licence including a microchip
Approval No: ...

1. Manufacturing brand or trademark: ...
2. Name of model: ...
3. Name of manufacturer or of its representative, where applicable:
4. Address of manufacturer or of its representative, where applicable:
5. Laboratory test reports:
6. Date of approval: ...
7. Date of withdrawal of approval: ...
8. Place: ...
9. Date: ...

10. Descriptive documents in Annex: ...

11. Signature: ...

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) No 383/2012. (See end of Document for details)

- (1) OJ L 403, 30.12.2006, p. 18.
- (2) OJ L 281, 23.11.1995, p. 31.

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