Status implementation 17025:2017 in the Netherlands

Presentation for workshop EPPO
September 2019

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Implementation 17025:2017

- Documents
- Process
- Result up to August 2019
- Main categories non conformities (NC's)

Documents

- Standard EN ISO/IEC 17025:2017
- RvA- T049 “Implementation of EN ISO/IEC 17025:2017”
  - Explanatory document
  - T-documents are mandatory
  - Describes the procedure and policy of the RvA concerning the transition to EN ISO/IEC 17025:2017

RvA- T049

- Assessments against EN ISO/IEC 17025:2017
- The decision process for accreditation against this new standard
- The replacement of the declarations and scopes of accreditations
- A comparison between EN ISO/IEC 17025:2005 and EN ISO/IEC 17025:2017

Process 1-2

- From January 2019, the regular assessments have been carried out against the requirements in EN ISO/IEC 17025:2017
- We make a distinction between NC's regarding 17025:2005 and 17025:2017
- For regular assessments taking place before 30 April 2020, the NC's regarding 17025:2017 can be assessed during that assessment

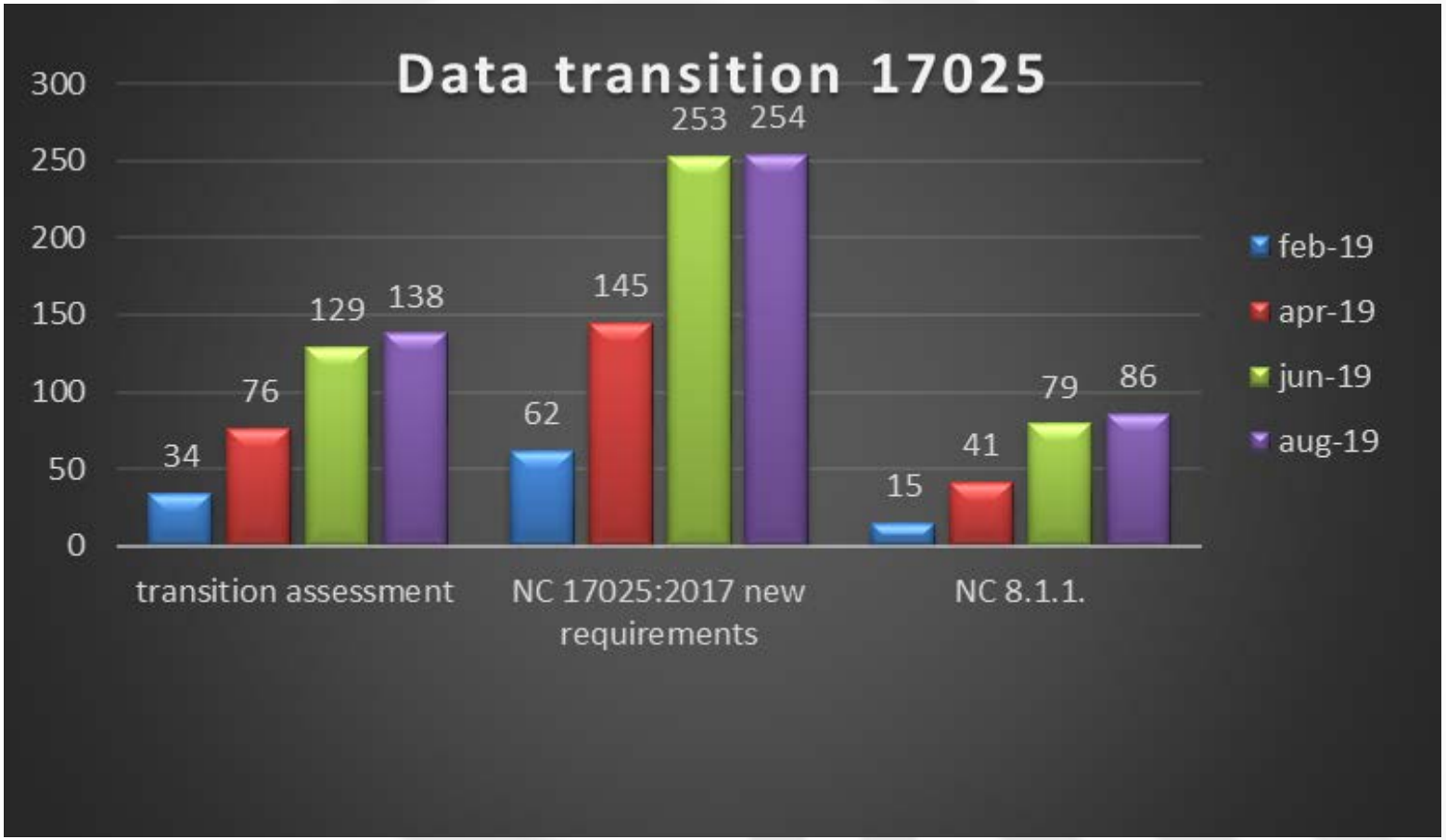
Process 2-2

- The organisation can implement corrective actions and submit its report before 1 September 2020
- Laboratories can request an additional assessment
- Otherwise, they risk not being accredited for 17025:2017 by 30 November 2020
- Accreditation based on EN ISO/IEC 17025:2005 will be withdrawn with effect from 30 November 2020.

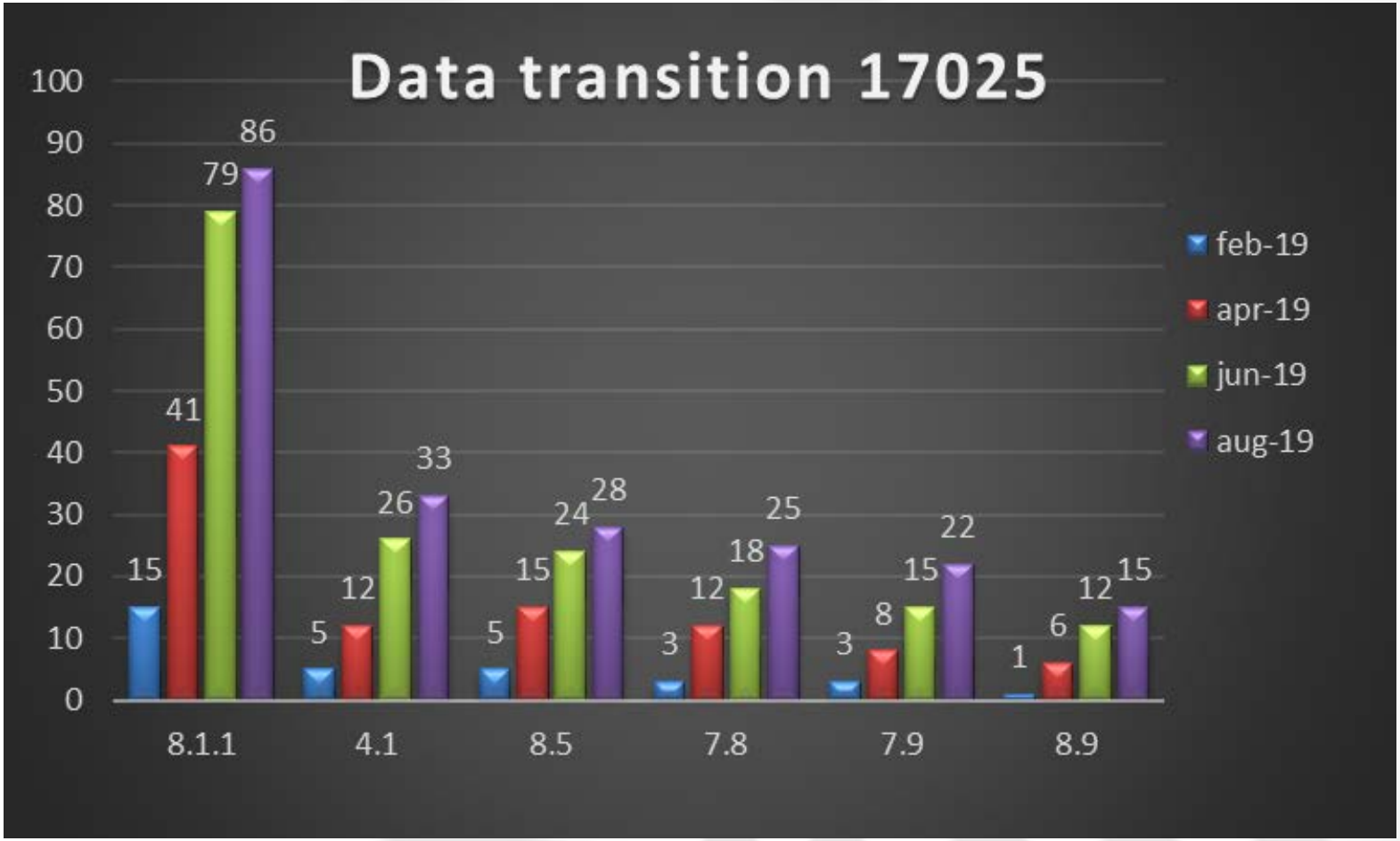
Result up to august 2019

- ± 300 accreditations 17025
- 138 assessments have been carried out against the 2017 version
- At the moment, 25 organisations have been accredited against against the 2017 version (6K and 19L)

Result up to august 2019



Main categories for non conformities



Conclusion

- Main NC's for 8.1.1
- Many laboratories have not yet fully completed the transition (8.1.1)
- They use 2019 for the transition
- other NC's for 4.1, 8.5, 7.8, 7.9 and 8.9