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|---|---|
| <b>87-1 (2004)</b>  | <b>Acetonitrile</b>   |
| CAS N°: 75-05-8   | EINECS N°: 200-835-2  |
| EC-LV (8 h): 70 mg/m<br>Lowest European LV (8h): 50 mg/m <sup>3</sup><br>Highest European LV (8h): 68 mg/m <sup>3</sup> | EC-STLV: -<br>Lowest European STLV: 100 mg/m <sup>3</sup><br>Highest European STLV: 280 mg/m <sup>3</sup> |

### SUMMARY OF THE METHOD

|                                      |  |
|--------------------------------------|--|
| <b>Language:</b><br>Spanish, English | <b>Reference:</b><br><b>Determinación de acetonitrilo en aire:</b> MTA/MA-055/A04, Métodos de Toma de muestra y análisis, INSHT (2004).<br><br><b>Acetonitrile</b> |
|--------------------------------------|--|

**Summary:** Air is drawn through a sorbent tube containing activated charcoal. After sampling, the collected acetonitrile is desorbed with dichloromethane containing methanol (50 µl/ml) and the resulting solution is analysed by GC/FID.

### SAMPLING

|                                  |                                |
|----------------------------------|--------------------------------|
| <b>Sampler type</b>              | Sorbent tube                   |
| <b>Sampling substrate</b>        | Activated charcoal (100/50 mg) |
| <b>Recommended flow rate</b>     | 0,2 l/min                      |
| <b>Recommended sampling time</b> | 20 min                         |
| <b>Recommended volume</b>        | 4 l                            |

### TRANSPORT AND STORAGE

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| <b>Description/conditions of transport and storage incl. specific issues</b> | Sample tubes are closed with plastic caps and transported to the laboratory as soon as possible. Stability was studied by sampling test gas atmosphere at RH = 80 %. It has shown that samples can be stored for up to 7 days at 4 °C (93,1 ± 1,83 % recovery). |
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### ANALYSIS

|                             |  |
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| <b>Sample preparation</b>   | The front and back sorbent sections are transferred to separate vials. The glass wool plugs are discarded. 1 ml of dichloromethane containing 50 µl of methanol and 1 µl of 1-hexanol as internal standard is added to each vial, which is then capped and allowed to stand for 30 min, with occasional agitation. |
| <b>Analytical technique</b> | Analysis by GC / FID.  |

### METHOD EVALUATION DATA

|                             |  |
|-----------------------------|--|
| <b>Range studied</b>        | 7 - 145 mg/m <sup>3</sup>                              |
| <b>Sampling bias</b>        | -  |
| <b>Analytical bias</b>      | -  |
| <b>Method bias</b>          | -6,1 % to +1,6 % (corrected for desorption efficiency) |
| <b>Sampling precision</b>   | -  |
| <b>Analytical precision</b> | 0,4 - 0,7 %  |
| <b>Method precision</b>     | 1 - 2,1 %  |

| METHOD EVALUATION DATA (continued)                               |  |
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| Limit of quantification  | -  |
| Overall uncertainty (EN 482)                                     | 3,7 - 8,6 %  |
| Expanded uncertainty (prEN 482)                                  | 13,2 - 14,8 %  |
| INFORMATION IN RELATION TO THE VALIDATION                        |  |
| Is the sample dissolution procedure described widely applicable? | yes  |
| Does the analysis include wall deposits, where applicable?       | not applicable   |
| Was a test gas atmosphere used, where applicable?                | Yes, dynamic test gas apparatus.   |
| How was the recovery determined?                                 | From spiked samples an analytical recovery of 100 % was found.<br>From test gas atmospheres studies a method recovery > 94 % was found.  |
| Was the sampler capacity or breakthrough volume determined?      | Yes. BV was determined to be 10 l sampling at a flow rate of 0,2 l/min, RH 83 %, an acetonitrile concentration of 125 mg/m <sup>3</sup> .  |
| Was temperature and RH considered, where appropriate?            | Yes. The method was evaluated by sampling generated atmospheres of acetonitrile at both high (80 %) and low (20 %) RH. The effect of temperature was not studied.  |
| EVALUATION   |  |
| Rating category  | A 1  |
| Rationale for rating   | Up to date methodology, detailed method description according ISO 78/2, performance and validation test data included in the method, OU and expanded uncertainty requirements met.<br><br>The overall uncertainty data above have been calculated from MTA/MA overall recovery and precision data using the formula in EN 482. The expanded uncertainty data have been calculated using the method described in the EU mandated report <i>Analytical methods for chemical agents</i> . |
| Observations   | -  |
| Similar methods  | None.  |