Appendix A: Guidelines for Measuring Methane and Nitrous Oxide Emissions from Rice Paddy Fields

This appendix explains how the methane (CH_4) and nitrous oxide (N_2O) emissions can be measured in rice paddy fields. It is necessary that the implementation of CH_4 and N_2O measurement by a closed chamber method involves technicians who have been authorized by the independent experts before the validation and operators who have been trained by the authorized technician.

In order to obtain the independent experts' advice on the competence of the technicians or to propose alternatives other than the ones explained in Appendix A to be included in this methodology with sufficient scientific evidence, firstly project participants need to contact the Ministry of Agriculture, Forestry and Fisheries of Japan as a representative of the Expert Committee established under the initiative of the Asian Development Bank for Climate-Resilient Agriculture and Low-Carbon Food Systems in the ASEAN Region as follows: maff JCMTML@maff.go.jp

The following tables are arranged sequentially from the chamber design to the calculation of the seasonal CH₄ and N₂O emissions and emission factors. See also "Guidelines for Measuring CH₄ and N₂O Emissions from Rice Paddies by a Manually Operated Closed Chamber Method" (pdf file, 8.4 MB) for the scientific basis and better understanding of the following guidance through the visual presentation.

Table A-1. Chamber design

| Feature | Conditions | | | |
|------------|--|--|--|--|
| Material, | In general, chamber shapes and materials are inseparable factors. In addition, | | | |
| color, and | chamber shapes allowed are dependent on a rice planting system | | | |
| shape | (transplanting or direct seeding). | | | |
| | Cylinder-shaped chambers with round basal area are usually made of | | | |
| | commercially available non-transparent plastic containers. Painting those with | | | |
| | whitish color, if not inherent, or a cover with reflective material is | | | |
| | recommended to prevent the increase in the inside temperature. Cylinder- | | | |
| | shaped chambers can be used only for the direct seeding system. | | | |
| | Rectangular-shaped chambers with square basal area are made of transparent | | | |

acrylic plates (with stainless steel frames for the reinforcement, if necessary). Rectangular-shaped chambers can be used for both transplanting and the direct seeding system.

The total chamber height (including that of a chamber base) is recommended to be higher than the rice plant height. Double- or triple-deck style is available to the rectangular-shaped chambers, which are adjustable depending on the growing plant height.

The constant placement of chambers equipped with upper lids that can be opened and closed is not recommended.

Base material and shape

The chamber base needs to be installed at least one day before the first gas sampling and must remain in the field throughout the season. Base materials and shapes depend on the chamber shapes. The aboveground height of the bases is recommended to be lower than or equal to 30 cm.

For cylinder-shaped chambers, a round-shaped base with a water sealing is usually made of plastic materials. A cylinder-shaped base in the soil requires holes on the sidewall to allow water exchange between inside and outside the chamber area.

For rectangular-shaped chambers, 4 corner pillars, made of PVC pipes or metal rods, stuck into the plow pan are sufficient when there is surface water. Top of the pillars are required to be underwater for sealing. When there is no or shallow surface water, chambers can be gently placed on the soil. CH₄ ebullition may happen after the placement, and it should be escaped from or well mixed in the chamber headspace before the first gas sampling. Square-shaped bases with water sealings made of plastic materials are also available temporarily or constantly during the season.

Basal area

The total area covered by chambers in one field is required to be wider than $0.25~\text{m}^2$. To accommodate this area, multiple ($n \ge 2$) chambers are used in one field. The minimum distance between each chamber should be 1 m. For the transplanting system, the basal area size of a rectangular-shaped chamber should be a multiple of rice plant density to appropriately capture GHG emissions (diffusion and ebullition) from the soil or the surface water. For example, if the plant density is 30 cm \times 15 cm, the basal area should be 30 cm

| | × 30 cm (covering 2 rice hills), 30 cm × 60 cm (4 hills), or etc. |
|-----------|--|
| Accessory | A chamber needs to be equipped with a gas sampling port, an inside fan, an inside thermometer, and an air vent. The port may also serve as the vent. The port should be apart from the chamber wall by using a tube connected to a stopcock. The dead volume in the tube should be replaced before the gas sampling. A weight, such as a battery for the fan operation, can be gently placed on the upper lid to prevent the chamber from falling down in windy weather. |
| | |

Table A-2. Gas sampling

| Feature | Conditions |
|---|--|
| Chamber area | A scaffold needs to be installed at least one day before the first gas sampling to reach the chamber areas without disturbing the soil. The chamber area needs to be apart 1.5 m from the ridge of the field. |
| Chamber replication per field | At least 2 chambers, depending on the basal area size (see the basal area feature in Table A-1). |
| Number of gas samples per chamber placement | At least 3 samples during the chamber closure time (30-40 min). The first gas sample should be collected after ≥1 min after the chamber placement to wait for the headspace gas to become well-mixed. |
| Gas sampling time of day | Morning, especially in the early hours (e.g., 7 am-10 am). If the sampling time must be extended to daytime, the schedule should be designed to prevent the systematic spatial bias since CH ₄ emissions are emitted more in daytime. |
| Frequency | At least once per week. To better trace the possible temporary CH ₄ emission peak during a drainage event and the possible temporary N ₂ O emission peak after nitrogen fertilizer topdressing, additional measurements once or twice are recommended during these events. |
| Gas | The gas sampled from the port should be stored into a glass or plastic evacuated |

| storage | vial (with a rubber stopper), a plastic or aluminum bag, or a plastic syringe. A gas leak test for the expected storage duration needs to be implemented before the start of the season and the gas concentration analyzed needs to be corrected appropriately, if applicable. |
|------------------|---|
| Manual operation | Uniform and gentle manual operation needs to be implemented regardless of time and place. Several operators should simultaneously implement the measurement in the reference fields and project fields. When moving the chamber from one location to another, the air inside the chamber should be replaced by pushing the chamber sideways and operating the fan. It is necessary to submit a film recording a series of gas sampling operations by the technicians to the independent experts for authorization of their skill prior to validation. If the technicians' skills are insufficient, the film shall be resubmitted in accordance with the independent experts' advice. |

Table A-3. Laboratory gas analysis

| Feature | Conditions |
|-----------|---|
| Method | A CH ₄ concentration needs to be analyzed by a gas chromatograph (GC) equipped with a flame ionization detector (FID) or a laser spectroscope. A N ₂ O concentration needs to be analyzed by a gas chromatograph (GC) |
| | equipped with an electron capture detector (ECD) or a laser spectroscope. In case of using a laser spectroscope ¹ , project participants should follow the manufacture's instruction for the gas analysis. |
| GC system | The GC system consists of a gas injection port, a separation column, a gas detector, a data processor, etc. Carrier gas and the standard gas are essential |
| | for the steady operation and the analysis. |
| | The ECD-GC should be equipped with a multi-port valve to remove oxygen and water vapor for the refined detection of N ₂ O. |
| | It is necessary to submit the column diagram and photos of GC appearance with and without opening the oven door to the independent experts before the |

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 $^{^{1}}$ A portable laser spectroscope connected online to multiple chambers is to be usable if the method is scientifically sound and if all the requirements are met.

| | validation. |
|---------------------------|--|
| Calibration line or curve | The certified standard gases need to be used to draw a calibration line or curve. 2-point calibration is sufficient for FID-GC using the CH ₄ standard gas with the atmospheric ambient concentration (e.g., \sim 2 ppm) and a higher concentration (e.g., 50-100 ppm). 2- or 3-point calibration is sufficient for ECD-GC using the N ₂ O standard gas with the atmospheric ambient concentration (e.g., \sim 0.3 ppm) and higher concentrations (e.g., 2-10 ppm). Note that the linearity is not always secured for ECD-GC to detect the higher concentration of N ₂ O. A calibration line or curve needs to be drawn each day before and after the |
| Quality | The repeatability of the GC analysis needs to be tested before the start of the season using the certified standard gases. The coefficient variation (CV) of $10\text{-}20$ repeated analyses of the same standard gas should be $\leq 5\%$ for all the used standard gases (i.e., ambient and higher concentrations). It is necessary to submit the results of the repeatability test to the independent experts for approval of quality control before the validation. If the results are poor (i.e., $CV > 5\%$), the result of additional blind test is recommended to submit. |

Table A-4. Calculation of the seasonal total emission of CH_4 or N_2O and emission factors

| Order | Procedure | | |
|-------|---|---|---|
| 1 | Calculate the mass of CH ₄ or N ₂ O in the analyzed gas sample: | | |
| | $m_t = c_t \times V \times M \times \frac{1atm}{R \times T_t \times 1000}$ | | |
| | Where: | | |
| | m_t | = | Mass of CH ₄ or N ₂ O in chamber at time t (mg) |
| | t | = | Time point of gas sampling (e.g. 1, 16, and 31 min after |
| | | | chamber placement in case of 3 samples for 30 min) |
| | c_t | = | $\mathrm{CH_4}$ or $\mathrm{N_2O}$ concentration in chamber at time t (ppm) |
| | V | = | Chamber volume (L) |
| | М | = | $Molar \ mass \ of \ CH_4 \ (16.042 \ g \ mol^{-1}) \ or \ N_2O \ (44.0128 \ g \ mol^{-1})$ |

| | 1atm | = | Assume constant pressure of 1 atm, unless the inside pressure |
|---|--|--|--|
| | | | is recorded |
| | R | = | Universal gas constant: 0.08206 L atm K ⁻¹ mol ⁻¹ |
| | T_t | = | Temperature at time $t(K)$ |
| 2 | Determine the slope of the line of best fit for the values of over time: $s = \frac{\Delta m}{\Delta t}$ | | |
| | | | |
| | Where: | Δt | |
| | S | = | Slope of line of best fit (mg min ⁻¹) |
| 3 | Calculate the hourly flux for one chamber measurement: | | |
| | | $F_{ch} = s \times \frac{\epsilon}{2}$ | 50min |
| | Where: | | А |
| | F_{ch} | = | Flux of chamber ch (mg m ⁻² h ⁻¹) |
| | ch | = | Index for replicated chamber in a field |
| | A | = | Chamber basal area (m²) |
| 4 | Calcula | te the averag | e hourly flux in a field: |
| | $F = \frac{\sum_{ch=1}^{n} F_{ch}}{n}$ | | |
| | Where: | n | |
| | F | = | Average flux of a field (mg m ⁻² h ⁻¹) |
| | n | = | Number of replicated chambers in a field |
| 5 | Calculate the total emission in one measurement interval: | | mission in one measurement interval: |
| | | $E_i = \frac{(F_i + F_i)}{2}$ | $(+1) \times 24h \times D_i$ |
| | Where: | v | 2 |
| | E_i | = | Total emission in interval $i \text{ (mg m}^{-2})$ |
| | i | = | Index for measurement interval in a season |
| | F_i | = | Hourly flux at the start of interval i (mg m ⁻² h ⁻¹) |
| | F_{i+1} | = | Hourly flux at the end of interval i (mg m ⁻² h ⁻¹) |
| | D_i | = | Number of days in interval i (d) |
| | Note the | at flux on pla | nting day and flux on harvest day can be assumed to be zero if |
| | measure | ement is not i | implemented on those days. |
| 6 | Calcula | te the season | al total emission in a field: |
| | | $E = \sum_{i=1}^{N} E_i$ | i |
| | 1 | | |

| | Where: | | |
|---|---|--|--|
| | Ε | = | Total emission in a season (mg m ⁻²) |
| | N | = | Number of measurement intervals in a season |
| 7 | Calculate the emission factor for the gas in stratum st in season s | | |
| | Where: | $EF_{s,st} = \frac{\sum_{f=1}^{F}}{}$ | $\frac{E_f \times 10^{-2}}{F}$ |
| | $EF_{s,st}$ | = | Emission factor of the gas in stratum st in season s (kg ha ⁻¹ season ⁻¹) |
| | E_f | = | Total emissions of the gas in field f of stratum st in season s |
| | п | | $(\text{mg m}^{-2} \text{ season}^{-1})$ |
| | F | = | Number of (representative) fields of stratum <i>st</i> in season <i>s</i> |
| 8 | Calculate the emission factor for the gas per day in stratum st in season s | | |
| | Where: | $EF_{s,d,st} = \frac{\sum_{f=1}^{F}}{1}$ | $\frac{1}{E_f \times 10^{-2}} \frac{E_f \times 10^{-2}}{D_f}$ |
| | $EF_{s,d,st}$ | = | Emission factor of the gas per day in stratum st in season s (kg |
| | | | $ha^{-1} day^{-1}$) |
| | E_f | = | Total emissions of the gas in field f of stratum st in season s |
| | | | $(\text{mg m}^{-2} \text{ season}^{-1})$ |
| | D_f | = | Total number of days in field f of stratum st in season s (days season ⁻¹) |
| | F | = | Number of (representative) fields of stratum <i>st</i> in season <i>s</i> |