Asynchrony in respiratory movements between the pulmonary lobes in patients with COPD: continuous measurement of lung density by 4-dimensional dynamic-ventilation CT

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Purpose: Four-dimensional dynamic-ventilation CT imaging demonstrates continuous movement of the lung. The aim of this study was to assess the correlation between interlobar synchrony in lung density and spirometric values in COPD patients and smokers, by measuring the continuous changes in lung density during respiration on the dynamic-ventilation CT.

Materials and methods: Thirty-two smokers, including ten with COPD, underwent dynamic-ventilation CT during free breathing. CT data were continuously reconstructed every 0.5 sec. Mean lung density (MLD) of the five lobes (right upper [RU], right middle [RM], right lower [RL], left upper [LU], and left lower [LL]) was continuously measured by commercially available software using a fixed volume of volume of interest which was placed and tracked on a single designated point in each lobe. Concordance between the MLD time curves of six pairs of lung lobes (RU-RL, RU-RM, RM-RL, LU-LL, RU-LU, and RL-LL lobes) was expressed by cross-correlation coefficients. The relationship between these cross-correlation coefficients and the forced expiratory volume in one second/forced vital capacity (FEV_{1.0}/FVC) values was assessed by Spearman rank correlation analysis.

Results: In all six pairs of the pulmonary lobes, the cross-correlation coefficients of the two MLD curves were significantly positively correlated with FEV_{1.0}/FVC ($\rho = 0.60-0.73$, $P < 0.001$). The mean value of the six coefficients strongly correlated with FEV_{1.0}/FVC ($\rho = 0.80$, $P < 0.0001$).

Conclusion: The synchrony of respiratory movements between the pulmonary lobes is limited or lost in patients with more severe airflow limitation.

Keywords: computed tomography, chronic obstructive pulmonary disease, emphysema, airflow limitation, mean lung density

Introduction

In patients with smoking-induced COPD, many previous publications analyzing radiological findings have reported heterogeneous distributions of emphysema, airflow, or even blood flow in the affected lung. For example, multiple papers have demonstrated that heterogeneous emphysema distribution and inhomogeneous cluster sizes of emphysema exist in the lung by analyzing chest CT,¹⁻⁵ that heterogeneous gas trapping and pulmonary volume changes are observed on paired inspiratory and expiratory CT scans;⁶⁻¹⁰ and that, on contrast-enhanced magnetic resonance (MR) imaging and hyperpolarized helium/xenon MR, both blood flow and airflow are distributed heterogeneously in the lung in COPD patients.¹¹⁻¹³ Similar inhomogeneous air

distribution in the lung by ventilation has also been confirmed by scintigraphy in COPD patients.¹⁴

While these observations of the COPD lung have been based on static imaging techniques, some reports using dynamic-ventilatory MR scanning have also demonstrated that asynchronized, heterogeneous movements are observed in the diaphragms or between the chest wall and the diaphragm.^{15–17} Although these MR scans are not able to visualize the motion of lung parenchyma directly, disconnected (paradoxical) respiratory movement between the chest wall and the diaphragm, which can be considered as the conventional “Hoover sign” at physical exam,¹⁸ implies that lung motion itself would be heterogeneous or asynchronized in patients with COPD.

Currently, 320-row and 256-row multidetector CT (MDCT) scanners can continuously scan the thorax under free breathing conditions, which first visualizes respiratory motion of the lung and bronchi directly and can be analyzed quantitatively using designated software or workstations.^{19,20} Although this 4-dimensional dynamic-ventilation CT currently scans a limited part of the whole lung (≤ 160 mm in length), the continuous mean lung density (MLD) values of the dynamic CT are measurable in the separated right and left lungs or at a certain tracked point in the lung using various software. Based on strong correlations between the MLD and lung volume (LV) values,^{21,22} the continuous change in MLD can be considered a good index to reflect respiratory changes in LV.

By using dynamic-ventilation CT, we have recently reported that asynchrony between lung movement and central airway movement exists in patients with severe COPD.²⁰ Although there is no published information on asynchrony or heterogeneity in respiratory movements between or among the lungs, based on previous observations, it can be predicted that dynamic-ventilation CT during respiration would demonstrate some heterogeneous lung movements in paired continuous MLD measurements and bring novel perspectives to better understand COPD physiology.

Thus, we aimed to investigate asynchrony/heterogeneity in continuous lung density measurement of the lung on dynamic-ventilation CT and to assess the correlation with spirometric values in smokers with and without COPD.

Materials and methods

This study was approved by the Institutional Review Board at Ohara General Hospital. Written informed consent from enrolled patients was waived, since all examinations in this study were performed as part of routine clinical care.

All patients' data including CT scans and spirometric values were anonymized for analyses.

Subjects

Thirty-two smokers (16 ex-smokers and 16 current smokers; four females and 28 males; mean age 70 ± 12 years) underwent spirometry and a chest CT at Ohara Medical Center attached to Ohara General Hospital. Among the 32 subjects, the data of 21 subjects were previously analyzed with a different objective for other research.²⁰ They were scanned for preoperative assessment of resectable thoracic neoplasms ($n=22$) or for an assessment of COPD according to the criteria of the American Thoracic Society and European Respiratory Society ($n=10$).²³ No patients had thoracic tumors with invasion of the chest wall or proximal airways.

CT scan

All patients were scanned on a 320-row MDCT scanner (Aquilion ONE, Toshiba Medical Systems, Otawara, Tochigi, Japan) for both conventional (static) and dynamic-ventilatory scans. Using the wide volume mode (non-helical mode), dynamic scanning was performed at a fixed point without bed movement, resulting in fluoroscopic images of 160 mm in length.^{19,20} Scanning field of view (FOV) was selected from two settings based on patient body habitus: 320 (medium, $n=14$) or 400 mm (large, $n=18$). Other scanning and reconstruction parameters for the dynamic-ventilation CT were as follows: tube currents = 40 mA; tube voltage = 120 kVp; rotation time = 0.35 sec; total scanning time = 4.0–6.5 sec; imaging FOV: 320 mm; collimation = 0.5 mm; slice thickness = 1 mm (without image interval or overlapping); reconstruction kernel = FC15 (for mediastinum); reconstruction interval = 0.5 sec/frame (total 8–13 frames); and reconstruction method = half reconstruction. Scan data were converted to CT images using an iterative reconstruction method (Adaptive Iterative Dose Reduction using Three Dimensional Processing [AIDR3D], mild setting). Before the dynamic-ventilatory scanning, patients were coached to repeat deep breathing. Radiologic technologists monitored patients' respiratory movements and confirmed that expiratory movement from the peak inspiratory phase to the peak expiratory phase was included during the scanning.

Conventional static chest CT was also performed using helical scanning to image the whole thorax. The parameters for the conventional chest CT were as follows: tube currents = automatic exposure control; tube voltage = 120 kVp; scanning method = helical scanning; rotation time = 0.35 sec; beam pitch = 0.828; imaging

FOV: 320 mm; collimation = 0.5 mm × 80; slice thickness = 1 mm; reconstruction kernel = FC17 (for mediastinum); iterative reconstruction = AIDR3D (mild setting).

Image analysis – continuous lung density measurement of the right and left lungs

Using commercially available software (Lung Volume Measurement, Toshiba Medical Systems), the MLD was measured automatically in each frame, and the time curve of the MLD on the dynamic-ventilatory scans was created. On the MLD curve, the inspiratory MLD was defined as the lowest MLD and expiratory MLD as the highest.

Since the whole lung could not be scanned within the 160 mm scan length, the LV was not measured in this experiment. The close correlation between changes in the MLD and those of LV was reported in a previous publication.²¹

Image analysis – continuous lung density measurement of the five lung lobes

Using another commercially available software program (PhyZiodynamics, Ziosoft Inc., Minato-ku, Tokyo, Japan), the MLD of each of the five lobes (right upper lobe [RUL], right middle lobe [RML], right lower lobe [RLL], left upper lobe [LUL], and left lower lobe [LLL]) was continuously measured. The software tracked on a single designated (center) point of a 3-dimensional spherical volume of interest (VOI) with a fixed volume (10 mm in radius) throughout the dynamic-ventilation CT (Figure 1, [Video S1](#)). Since the dynamic CT did not scan the whole thorax, the VOIs

Figure 1 Example of continuous mean lung density measurements in the right upper lobe using PhyZiodynamics software (Ziosoft Inc., Minato-ku, Tokyo, Japan).

Note: A spherical volume of interest (VOI) with a fixed volume (10 mm in radius) is placed and the center point of the VOI is tracked throughout the dynamic computed tomography (also see Supplementary materials).

Table 1 Clinical characteristics of 32 subjects

	Mean ± SD	(Range)
Gender (female:male)	(4:28)	–
Age (years)	70±12	(36 to 84)
Smoking index (pack-years)	46±21	(2 to 100)
Smoking status	(16:16)	–
(ex-smoker:current smoker)		
FEV _{1.0} /FVC	0.73±0.12	(0.41 to 0.92)

Abbreviation: FEV_{1.0}/FVC, forced expiratory volume in one second/forced vital capacity.

were placed at the following points in the five lung lobes: S3 in the RUL and LUL; S4 in the RML; S6 in the RLL and LLL.

Similar to the right and left lung measurements, the inspiratory MLD was defined as the lowest MLD and expiratory MLD as the highest on the MLD curve in each lobe.

Image analysis – emphysema measurement on conventional chest CT

On the conventional helical CT scans, an emphysema measurement was performed using the LV measurement software. The percentage low attenuation volume (LAV%, <−950 HU) of the whole lung was obtained automatically.

Spirometry

All subjects performed spirometry, including forced expiratory volume in one second (FEV_{1.0}) and forced vital capacity (FVC), according to American Thoracic Society standards.²³ The spirometric values from the study participants are shown in Table 1. Spirometry was performed within 2 weeks of the chest CT.

Statistical analysis

Cross-correlation coefficients ($-1 \leq R \leq 1$) were calculated between the two time curves of the MLD (right–left lungs and six pairs of lung lobes). If the two time curves have similar shapes in direction over time, the coefficient approaches one.^{19,20} Spearman rank correlation analysis was used to evaluate the associations between the CT indices and the FEV_{1.0}/FVC.

A *P*-value of <0.05 was considered significant. All statistical analyses were performed using JMP 12.0 software (SAS Institute Inc., Cary, NC, USA).

Results

Radiation dose assessment for dynamic-ventilation CT

For a single gantry rotation of 160 mm (0.35 sec), the CT dose index volume was 1.28 mGy for the medium scanning

FOV or 1.41 mGy for the large FOV. The dose-length product value for a single rotation was 20.4 mGy cm (medium FOV) or 22.5 mGy cm (large FOV). The total estimated radiation exposure for the dynamic-ventilation CT for 4.0–6.5 sec varied from 3.7 to 6.1 mSv (mean 5.4 mSv).

Dynamic MLD and static LAV% measurements

During the dynamic-ventilation CT, the change in MLD ranged from 46.5 to 84.6 HU in the right/left lungs and in each lung lobe (right lung, 65.6 HU; left lung, 67.6 HU; RUL, 47.6 HU; RML, 46.5 HU; RLL, 84.6 HU; LUL, 52.2 HU; LLL, 78.3 HU, respectively). The changes in MLD were larger in the lower lobes than in the upper lobes. All MLD values, particularly at expiratory phase, were significantly correlated with $FEV_{1.0}/FVC$ (Table 2, [Video S2](#)). The expiratory MLD of the right and left lungs achieved the highest and the second highest correlation coefficients to $FEV_{1.0}/FVC$ (the right lung, $\rho=0.78$; the left lung, $\rho=0.74$; $P<0.0001$ for both). The LAV% of the whole lung also significantly correlated with $FEV_{1.0}/FVC$ ($\rho=-0.73$, $P<0.0001$) (Table 2).

Synchrony of MLD curves between the right and left lungs

The cross-correlation coefficient between the two MLD curves of the right and left lungs was significantly correlated

with the $FEV_{1.0}/FVC$ ($\rho=0.66$, $P<0.0001$) (Table 3, Figure 2). This implies that the synchrony between the right and left lung movements was limited in severe COPD patients compared with smokers without airflow limitation.

Synchrony of MLD curves between the paired lung lobes

Similar to the cross-correlation between the right and left lungs, the cross-correlation coefficients of the two MLD curves from all six pairs of the pulmonary lobes were significantly positively correlated with $FEV_{1.0}/FVC$ ($\rho=0.60-0.73$, $P<0.001$) (Table 3, Figure 3). Further, the mean value of the six coefficients from the six lobar pairs strongly correlated with $FEV_{1.0}/FVC$ ($\rho=0.80$, $P<0.0001$) (Figure 4), which was the highest correlation in this study. These observations suggested that the synchrony of respiratory movements among the lung lobes was limited and lost in severe COPD patients.

Correlation between synchrony of MLD curves and emphysema index

Significant and insignificant correlations were found between LAV% of the whole lung and cross-correlation coefficients of paired MLD curves (Table 4). This implies that the extent of emphysema may partially cause asynchrony of inter-lung and interlobar movements but that emphysema severity would

Table 2 Measurements of lung density and correlations to $FEV_{1.0}/FVC$

	Density measurement		Correlation to $FEV_{1.0}/FVC$	
	Mean $\pm$ SD	(Range)	Coefficient (ρ)	P-value
Dynamic-ventilation CT				
Inspiratory-MLD (HU)				
Right lung	-874.9 $\pm$ 38.1	(-941.7 to -814.8)	0.62	<0.001
Left lung	-866.9 $\pm$ 38.6	(-953.5 to -809.9)	0.45	<0.01
RUL	-909.3 $\pm$ 34.3	(-959.8 to -836.2)	0.52	<0.01
RML	-908.8 $\pm$ 38.1	(-987.5 to -828.1)	0.44	<0.05
RLL	-861.6 $\pm$ 58.0	(-969.5 to -728.0)	0.49	<0.01
LUL	-907.8 $\pm$ 41.9	(-1,021.4 to -818.2)	0.51	<0.01
LLL	-866.9 $\pm$ 62.4	(-971.8 to -733.5)	0.42	<0.05
Expiratory-MLD (HU)				
Right lung	-809.4 $\pm$ 57.6	(-937.6 to -704.7)	0.78	<0.0001
Left lung	-799.4 $\pm$ 60.5	(-917.6 to -684.3)	0.74	<0.0001
RUL	-861.7 $\pm$ 49.3	(-946.8 to -751.5)	0.70	<0.0001
RML	-862.2 $\pm$ 47.7	(-945.9 to -765.5)	0.59	<0.001
RLL	-777.1 $\pm$ 86.6	(-936.2 to -625.1)	0.56	<0.001
LUL	-855.6 $\pm$ 61.6	(-1,003.4 to -731.9)	0.67	<0.0001
LLL	-788.5 $\pm$ 86.9	(-960.2 to -641.4)	0.60	<0.001
Helical CT				
LAV% (%)				
Whole lung	10.8 $\pm$ 14.8	(0 to 52.0)	-0.73	<0.0001

Abbreviations: $FEV_{1.0}/FVC$, forced expiratory volume in one second/forced vital capacity; LAV%, percentage low attenuation volume (<-950 HU); LLL, left lower lobe; LUL, left upper lobe; MLD, mean lung density; RLL, right lower lobe; RML, right middle lobe; RUL, right upper lobe.

Table 3 Cross-correlation coefficients between two MLD curves of the right and left lungs and those of paired lung lobes and their correlations to $FEV_{1.0}/FVC$

Pairs – MLD curves	Cross-correlation coefficient		Correlation to $FEV_{1.0}/FVC$	
	Mean $\pm$ SD	(Range)	Coefficient (ρ)	P-value
Right – left lungs	0.954 $\pm$ 0.049	(0.798 to 0.996)	0.66	<0.0001
RUL – RLL	0.784 $\pm$ 0.366	(–0.791 to 0.993)	0.67	<0.0001
RUL – RML	0.823 $\pm$ 0.428	(–0.984 to 0.999)	0.61	<0.001
RML – RLL	0.707 $\pm$ 0.495	(–0.968 to 0.996)	0.72	<0.0001
LUL – LLL	0.842 $\pm$ 0.298	(–0.460 to 0.997)	0.60	<0.001
RUL – LUL	0.928 $\pm$ 0.160	(0.183 to 0.999)	0.60	<0.001
RLL – LLL	0.801 $\pm$ 0.324	(–0.610 to 0.998)	0.73	<0.0001
Average of six lobar pairs	0.814 $\pm$ 0.259	(0.065 to 0.994)	0.80	<0.0001

Abbreviations: $FEV_{1.0}/FVC$, forced expiratory volume in one second/forced vital capacity; LLL, left lower lobe; LUL, left upper lobe; MLD, mean lung density; RLL, right lower lobe; RML, right middle lobe; RUL, right upper lobe.

not be the sole factor of heterogeneous lung movements observed in severe COPD patients.

Discussion

In this study, we first examined continuous MLD measurement on dynamic-ventilation CT in the separated right and left lungs and in five lung lobes and found that synchrony in MLD curves between the paired right-left lungs or paired lung lobes was reduced in severe COPD patients. These observations lead to the concept that respiratory movement of the lung would be heterogeneous or asynchronized (pantomime-like movement) as COPD progresses. The dynamic-ventilation CT which directly depicts respiratory lung movement shows the previously unknown, ambiguous physiological phenomena of COPD, which may be expanded to therapeutic strategies for the management of COPD.

Although this is the first study demonstrating heterogeneous asynchronized lung movement using 4-dimensional

CT, inhomogeneous air distribution by ventilation in COPD patients has been demonstrated by several different imaging modalities, including xenon-inhaled (xenon-enhanced) CT scanning using a dual-energy technique.^{24–26} The latest study analyzing xenon-CT in COPD patients has shown that decreased air distribution is observed even in the non-emphysematous, apparently normal lung areas, suggesting the presence of other coexisting conditions such as proximal or small airway disease in COPD.²⁵ Considering that inhomogeneous air distribution by ventilation has also been demonstrated by paired inspiratory and expiratory CT^{6–9} and hyperpolarized xenon or helium MR,^{12,13} our observation of asynchronized, heterogeneous lung movement by ventilation in COPD is consistent with these previous publications and further expands the 4-dimensional imaging approach.

It is still difficult to explain why this heterogeneous, asynchronized respiratory movement in the lung is observed on dynamic-ventilation CT in patients with COPD. However,

Figure 2 Mean lung density (MLD) time curves of the right and left lungs.

Notes: In a smoker without COPD (**A**), the two MLD curves are comparable, suggesting that both lungs move simultaneously (cross-correlation coefficient [CCC] = 0.982). In contrast, in a COPD patient (**B**), the right and left MLD curves are uneven and not well-matched, suggesting the occurrence of asynchronous lung movements (CCC = 0.798).

Abbreviations: $FEV_{1.0}/FVC$, forced expiratory volume in one second/forced vital capacity; LAV%, percentage low attenuation volume (<–950 HU).

Figure 3 Mean lung density (MLD) time curves of three lung lobes in the right lung.

Notes: In a smoker without COPD (**A**), the three MLD time curves of the three lung lobes have similar shapes, which indicates synchronous respiratory movements of the lobes (cross-correlation coefficient [CCC] of three pairs =0.975–0.996). A larger change in MLD in the lower lobe than in the upper and middle lobes suggests that respiratory volume changes mainly occur in the lower lobe in this subject. In a severe COPD patient (**B**), the three MLD time curves are unmatched, resulting in uncertain inspiratory/expiratory movements. Note that nearly opposite MLD curves of the middle and lower lobes are observed (CCC =−0.135), suggesting that asynchronized respiratory airflow exists in this patient.

Abbreviations: FEV_{1.0}/FVC, forced expiratory volume in one second/forced vital capacity; LAV%, percentage low attenuation volume (<−950 HU); RUL, right upper lobe; RML, right middle lobe; RLL, right lower lobe.

there are several potential components that prevent uniform (homogeneous) lung movement in COPD. For example, inhomogeneous distribution and heterogeneous progress of pulmonary emphysema must cause heterogeneous respiratory movement of the lung; more progressed, emphysematous parts would not move largely during ventilation, while the normal or less affected lung areas would greatly inflate/deflate. Also, various degrees of proximal airway disease at different points in COPD can lead to heterogeneous respiratory movement of the lung ([Video S2](#)). The pulmonary lobe or segment with more severe airway

disease (bronchial wall thickening and narrowed luminal size) would show delayed, prolonged expiration compared with other lobes/segments without severe airway disease. Further, disconnection between chest wall movement and diaphragmatic movement and those between the right and left diaphragmatic movement in COPD, which has already been reported by dynamic-ventilation MR,¹⁶ are also important to understand heterogeneous lung movements between the right and left lungs and between the upper and lower lobes observed in this study. In other words, all significant factors of COPD, such as emphysema, airway disease, and abnormalities of extrapulmonary structures (flattened diaphragms or thinned respiratory muscles), may

Figure 4 Correlation between spirometric value (FEV_{1.0}/FVC) and mean cross-correlation coefficient (CCC) from six pairs of lung lobes.

Note: Most smokers without COPD have CCC larger than 0.9, suggesting that respiratory movement is uniform and synchronized among lung lobes in non-COPD smokers.

Abbreviation: FEV_{1.0}/FVC, forced expiratory volume in one second/forced vital capacity.

Table 4 Correlation between cross-correlation coefficient of paired MLD curves and emphysema index of the whole lung

Pairs – MLD curves	Correlation to LAV% (whole lung)	
	Coefficient (p)	P-value
Right – left lungs	−0.44	<0.05
RUL – RLL	−0.40	<0.05
RUL – RML	−0.63	<0.001
RML – RLL	−0.54	<0.01
LUL – LLL	−0.31	NS
RUL – LUL	−0.52	<0.01
RLL – LLL	−0.43	<0.05
Average of six lobar pairs	−0.54	<0.01

Abbreviations: LAV%, percentage low attenuation volume (<−950 HU); LLL, left lower lobe; LUL, left upper lobe; MLD, mean lung density; NS, not significant; RLL, right lower lobe; RML, right middle lobe; RUL, right upper lobe.

integratively lead to asynchronized, heterogeneous lung movements in COPD.

If asynchrony or heterogeneity in lung movement is truly important to understand COPD physiology, it would be of interest to monitor the change in the asynchrony/heterogeneity before and after therapeutic interventions in COPD patients. Medications such as bronchodilators and also respiratory rehabilitation methods may result in a decrease in the asynchrony/heterogeneity of lung movement. Suga et al have reported that asynchronized, paradoxical movements of the right and left diaphragms are restored after LV reduction surgery, as evaluated with dynamic-ventilation MR.¹⁷ We have started planning future studies in order to expand our observations to clinical care of COPD patients, based on a prediction that the asynchrony/heterogeneity of lung movement can be used as a comprehensive physiological CT index in COPD and would be improved after a combination of different therapeutic approaches for COPD patients. Also, MLD curves from the two adjacent lobes with very different or opposite shapes (Figure 3) may imply the presence of collateral ventilation between these lobes. Since collateral ventilation prevents successful endobronchial valve treatment for severe COPD patients,²⁷ dynamic-ventilation CT may have a clinical role, similar to the conventional catheter-based endobronchial measurement method, in determining the usefulness of valve treatment. Although this study demonstrates purely physiological aspects of COPD by using dynamic-ventilation CT, future studies may reveal more advantages of dynamic-ventilation CT in daily clinical care for COPD patients, such as the combination of paradoxical airway movement²⁰ and heterogeneity of lobar movement which may be a target for intrabronchial stenting or other therapeutic options.

This study has several limitations which should be noted. First, the total number of enrolled subjects was small, particularly the number of COPD patients. The very small number of female COPD patients (n=2) may raise the question of whether our observations are applicable to all COPD patients. Also, many enrolled patients underwent the dynamic CT scan as a preoperative assessment for tumor invasion or adhesion to the pleura. Although we confirmed that no patient had a large tumor which invaded the adjacent structures or occluded the proximal bronchi, our observations should be reproduced among smokers/COPD patients without thoracic neoplasms in the future. Second, the dynamic-ventilation CT could not scan the entire thorax. Although we believed that it was best at the moment to utilize the MLD measurement instead of right and left LV measurements, true respiratory changes in LV

could not be measured. Third, due to the limited scan field, the VOI in each lung lobe was small and it was unclear whether or not the measured MLD in the VOI truly represented the respiratory movement of the targeted lobe. Since we found that interlobar asynchrony or heterogeneity in respiratory movement existed in severe COPD patients, such asynchrony may be observed even in the single lung lobe, which was not able to be assessed by our current method. In future studies, the current technique should be improved by multiple VOI measurements in each lobe or more comprehensive, dynamic measurements of the entire scanned lung. Also, weaker correlations between spirometric values and expiratory MLD of the lower lobes than those of the upper lobes, which is discordant to previous studies analyzing CT-based LV on combined inspiratory and expiratory scans,⁹ may have been due to the small VOI which could not cover the entire lower lobes. Fourth, we evaluated correlations between CT data and spirometric values only, which should be expanded with more physiological and clinical parameters such as plethysmographic values or the 6-minute walk test.

Conclusion

We have reported that the synchrony of respiratory movements between the right and left lungs and between the pulmonary lobes is limited or lost in COPD patients with more severe airflow limitation. Asynchrony/heterogeneity in lung movements, which can be exclusively observed by the 4-dimensional dynamic-ventilation CT, may be an important clue to comprehend the pathophysiology of COPD.

Acknowledgments

University of the Ryukyus, Ohara General Hospital, and Shiga University of Medical Science received a research grant from Toshiba Medical Systems. University of the Ryukyus also received a research grant from Ziosoft Inc.

This retrospective study was arranged as part of the Area-detector Computed Tomography for the Investigation of Thoracic Diseases (ACTive), a multicenter research alliance in Japan. The authors thank Mr Shinsuke Tsukagoshi, Mr Tatsuya Kimoto, (Toshiba Medical Systems), and Mr Yasuhiro Kondo (Ziosoft) for their technical support.

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Disclosure

Dr Yamashiro receives a research grant from the Japan Society for the Promotion of Science (Kakenhi-16K19837). The authors report no conflicts of interest in this work.

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