

CYCLE ANALYSIS OF GAS TURBINE-FUEL CELL  
HYBRID MICRO GENERATION SYSTEM

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**ABSTRACT**

Small distributed generation systems are currently attracting much attention because of their high energy utilization efficiency. Among them, a hybrid system based on micro gas turbine ( $\mu$ GT) and solid oxide fuel cell (SOFC) is expected to achieve much higher efficiency than a traditional  $\mu$ GT. In this paper, we investigate the effects of cycle design parameters on the performance and feasibility of a  $\mu$ GT-SOFC hybrid system for small apartments and businesses. As a result, a general design strategy is found that less direct fuel input to combustor as well as higher recuperator effectiveness leads to higher generation efficiency, while higher steam-carbon ratio moderates requirements for the material strength. It is also confirmed that the hybrid system is much superior to the recuperated gas turbine in terms of its power efficiency and aptitude for small distributed generation. A conceptual design of a 30kW  $\mu$ GT-SOFC hybrid system, of which diameter and height are 750mm and 1500mm, respectively, is shown to give power efficiency over 65% (LHV) in the best possible case.

**NOMENCLATURE**

**Symbols**

$F$ : Farady constant	C/mol
$G$ : Gibbs free energy	J/kg
$H$ : enthalpy	W
$J$ : cell current density	A/ m <sup>2</sup>
$K$ : equilibrium constant	
$Q$ : heat	W

$R$ : gas constant	J/mol K
$T$ : temperature	K
$V$ : voltage	V
$W$ : power	W
$i$ : current	A
$n$ : number of moles	mol
$p$ : pressure	atm
$x$ : mole fraction	
$\delta$ : thickness	mm
$\eta$ : efficiency	
$\rho$ : electrical resistance	$\Omega$ cm

**Subscripts**

cell	: SOFC cell
ref	: reforming or reformer
shf	: shifing
an	: anode
ca	: cathode
el	: electrolyte
net	: net value
act	: activation porarization
SOFC	: solid oxide fuel cell
c	: compressor
t	: turbine
DA	: inverter
gen	: generater

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### 3. CYCLE ANALYSIS

#### 3.1 Assumptions

The basic conditions of the present cycle analysis are shown in Table 1. Unless otherwise stated, these conditions are assumed. We make the following assumptions:

- (1) Gases do not leak outside the system.
- (2) Heat loss is negligibly small.
- (3) Chemical reactions proceed to equilibrium states.
- (4) Internal distributions of the temperature, gas composition, and pressure in each system component are not considered.
- (5) Steady state operation is considered.
- (6) Heat required in the reformer is supplied from the cell.
- (7) The anode outlet temperature, the cathode outlet temperature, and the reformer outlet temperature are all equal to the cell temperature.
- (8) In the combustor, the residual gas from the anode and the injected methane are burnt completely.

#### 3.2 Reaction Heats and Equilibrium Constants

For analyzing the reaction in the reformer and the cell, the reaction heats and the equilibrium constants should be known. The reaction heats are obtained from the data of the formation enthalpy in JANAF table <sup>(5)</sup>, except the formation enthalpy of methane, of which coefficient is given in the literature <sup>(6)</sup>. The equilibrium constants are obtained from difference of the Gibbs energies using the following equation:

$$K = \text{Exp}\left(-\frac{\Delta G}{RT}\right) \quad (1)$$

The reaction heats and equilibrium constants are expressed as polynomials fitting functions of temperature.

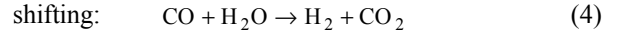
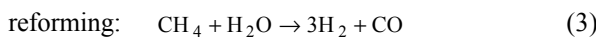
#### 3.3 Micro Gas Turbine

For the calculation of compression and expansion processes in the compressor and turbine, the standard method in the literature <sup>(7)</sup> is used. The combustor outlet temperature is first assumed, then the combustor methane input is determined by iterative calculation. The power output from the gas turbine is obtained by using the following equation:

$$W_{gt} = \eta_{gen} (\eta_{mgt} W_t - W_c) - W_{gcgt} \quad (2)$$

#### 3.4 Reformer

In the reformer, hydrogen and carbon oxide are produced from methane and steam. Inside the reformer, the reactions bellow are considered.



Reforming is an endothermic reaction, where shifting is an exothermic reaction. As a whole, the reaction in the reformer is endothermic. To allow methane and steam to react, the residual heat in the cell is transferred to the reformer.

The equilibrium constant of reforming  $K_{ref}$  and that of shifting  $K_{shf}$  are defined based on the partial pressures of species as:

$$K_{ref} = \frac{p_{\text{H}_2}^3 p_{\text{CO}}}{p_{\text{CH}_4} p_{\text{H}_2\text{O}}} \quad (5)$$

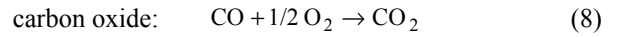
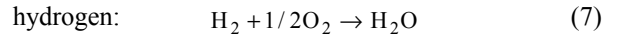
$$K_{shf} = \frac{p_{\text{H}_2} p_{\text{CO}_2}}{p_{\text{CO}} p_{\text{H}_2\text{O}}} \quad (6)$$

We can obtain mole numbers for reforming  $\Delta n_{ref}$  and for shifting  $\Delta n_{shf}$  by iterative calculation from  $K_{ref}$  and  $K_{shf}$  obtained using the method in 3.2. To estimate  $K_{ref}$  and  $K_{shf}$  we assume that the reformer exit temperature equals the cell temperature, and the reforming temperature equals the arithmetic mean of inlet and outlet temperatures. Once the mole numbers to react are estimated, the reaction heat  $Q_{ref}$  can be attained from overall heat balance.

### 3.5 Fuel Cell

#### 3.5.1 Theoretical voltage

In the cell, hydrogen and carbon oxide reacts and the electric power is generated through the following process:



with  $\Delta G_{\text{H}_2}$  representing the change of Gibbs energy before and after the reaction of hydrogen, the theoretical voltage is obtained as follows <sup>(8)</sup>:

$$V_{\text{H}_2} = -\frac{1}{2F} \Delta G_{\text{H}_2} - \frac{RT}{2F} \ln \left( \frac{p_{\text{an},\text{H}_2\text{O}}}{p_{\text{an},\text{H}_2} \sqrt{p_{\text{ca},\text{O}_2}}} \right) \quad (9)$$

Using equation (1), we can obtain:

$$V_{\text{H}_2} = \frac{RT}{2F} \left\{ \ln K_{\text{H}_2} - \ln \left( \frac{x_{\text{an},\text{H}_2\text{O}}}{x_{\text{an},\text{H}_2}} \right) + \frac{1}{2} \ln (x_{\text{ca},\text{O}_2} p_{\text{ca}}) \right\} \quad (10)$$

Similarly, for carbon oxide:

$$V_{\text{CO}} = \frac{RT}{2F} \left\{ \ln K_{\text{CO}} - \ln \left( \frac{x_{\text{an},\text{CO}_2}}{x_{\text{an},\text{CO}}} \right) + \frac{1}{2} \ln (x_{\text{ca},\text{O}_2} p_{\text{ca}}) \right\} \quad (11)$$

The equilibrium constants are obtained as described in 3.2.

#### 3.5.2 Reacting species

From the definition of equilibrium constants, we can obtain the oxygen partial pressure in the anode as follows:

$$p_{\text{an},\text{O}_2} = \left( \frac{1}{K_{\text{H}_2}} \frac{p_{\text{an},\text{H}_2\text{O}}}{p_{\text{an},\text{H}_2}} \right)^2 = \left( \frac{1}{K_{\text{CO}}} \frac{p_{\text{an},\text{CO}_2}}{p_{\text{an},\text{CO}}} \right)^2 \quad (12)$$

Using equation (12), we can confirm that the theoretical voltage of hydrogen (equation (10)) and that of carbon oxide (equation (11)) are equal.

At first, the amount of hydrogen to react is calculated. Then,

we can obtain the amount of carbon oxide to react using equation (12). Using the arithmetic mean of mole numbers at the cell inlet and outlet, the relationship between the amounts of reacting hydrogen and carbon oxide is obtained as:

$$\frac{1}{K_{CO}} \frac{n_{an,CO_2} + 1/2\Delta n_{CO}}{n_{an,CO} - 1/2\Delta n_{CO}} = \frac{1}{K_{H_2}} \frac{n_{an,H_2O} + 1/2\Delta n_{H_2}}{n_{an,H_2} - 1/2\Delta n_{H_2}} \quad (13)$$

where  $\Delta n_{H_2}$  and  $\Delta n_{CO}$  represent the amounts of reacting hydrogen and carbon oxide, respectively. Introducing  $\Delta n_{H_2}$  to equation (13),  $\Delta n_{CO}$  is estimated.

### 3.5.3 Polarization

In a real cell, the net voltage is lower than the theoretical voltage because of some drops in voltage (polarization). There are three kinds of polarizations, namely, activation polarization, ohmic polarization, and concentration polarization.

The activation polarization is estimated as follows <sup>(8)</sup>:

$$V_{act} = \left\{ V_a \frac{\ln(J_c/J_o)}{\ln(1+J_c/J_o)} \right\} \ln(1+J/J_o) \quad (14)$$

where

$$V_a = 0.074 (T / 1273) \quad (15)$$

$$J_o = 10^{(-A/T+B)} \quad (16)$$

$$J_c = 0.4 \quad (17)$$

The ohmic polarization is estimated as follows <sup>(3)</sup>:

$$\rho_{an} = 0.00298 \text{ Exp}(-1392 / T) \quad (18)$$

$$\rho_{ca} = 0.008114 \text{ Exp}(600 / T) \quad (19)$$

$$\rho_{el} = 0.00294 \text{ Exp}(10350 / T) \quad (20)$$

The concentration polarization is negligibly small if compared to the other polarizations <sup>(8)</sup>.

### 3.5.4 SOFC output

Considering the activation and ohmic polarizations, the net voltage of the cell is calculated as:

$$V_{net} = V_{H_2} - V_{act} - J(\rho_{an} \delta_{an} + \rho_{ca} \delta_{ca} + \rho_{el} \delta_{el}) \quad (21)$$

The current is obtained from the amount of reaction species as:

$$i = 2F\Delta n_{H_2} + 2F\Delta n_{CO} \quad (22)$$

Therefore, the SOFC output is:

$$W'_{SOFC} = V_{net} i \quad (23)$$

Now, we should take into account the power for SOFC auxiliaries. Then, the SOFC net output is obtained as follows:

$$W_{SOFC} = \eta_{DA} W'_{SOFC} - W_{gfc} - W_{blow} \quad (24)$$

### 3.5.5 Cell energy balance

From the total inlet enthalpy  $H_{in}$  and the reaction enthalpy  $\Delta H_{cell}$ , the total cell exit enthalpy  $H_{out}$  is obtained as:

$$H_{out} = \{H_{in} + (-\Delta H_{cell})\} - (W'_{SOFC} + Q_{ref}) \quad (25)$$

Assuming that the anode and cathode outlet temperatures are equal, they are obtained from  $H_{out}$ . We can derive the fuel input to SOFC for a certain cell temperature by iterative calculation.

A part of the anode exit gas is recirculated. The amount of recirculated gas is obtained from the specified S/C ratio. In this study, the generation efficiency is defined based upon the sum of SOFC and gas turbine heat inputs, which are estimated in terms of the lower heating value.

## 4. CHARACTERISTICS OF HYBRID SYSTEM

The  $\mu$ GT-SOFC hybrid system is expected to achieve efficiency near 70%. To clarify the major reasons, we performed an exergy analysis (Fig. 2). TIT is increased to 1200°C using a ceramic turbine. Each exergy fraction is defined as the ratio of output or exergy loss to exergy of fuel input. Thus, the exergy fraction of output is not equal to the generation efficiency. The combustion loss of the hybrid is lower than that of traditional recuperated gas turbine, because electric power and heat are generated simultaneously. The exhaust loss and the recuperator loss are lower than that of atmospheric SOFC. With the turbine, additional power is recovered effectively from high temperature exhaust gas. Smaller difference between the high and low temperatures in the recuperator makes the irreversibility of heat transfer decrease.

Generally, smaller power output tends to decrease the generation efficiency. To estimate the effect of reduction in power level on generation efficiency, we performed a cycle analysis for 30kW and 300kW systems of both hybrid system and traditional recuperated gas turbine. We assume that the compressor, turbine and mechanical efficiencies depend on the output level. To give these elemental efficiencies, it is also assumed that approximately one fifth of the total output is gained by the gas turbine. The result is shown in table 2. For the hybrid system, the difference in the heat input per 1kW output between 30kW and 300kW systems is 0.07kW, where that of traditional recuperated gas turbine is 0.23kW. It is evident that the hybrid system efficiency is less dependent on the power output level, so that the hybrid system is more suitable for small systems.

The gas turbines generation efficiency is generally decreased when the intake air temperature is elevated. This tendency is a disadvantage for a small generation system because the demand for electricity is high in summer. We estimated the effect of atmosphere temperature on the generation efficiency. The cycle calculations for compressor inlet temperature of 15°C and 30°C were performed for both a hybrid system and a traditional gas turbine (Table 3). For the hybrid system, the difference of heat input per 1kW output at 15°C and 30°C equals 0.03kW, while that of the traditional recuperated gas turbine is 0.12kW. Thus the hybrid system deteriorates less in its efficiency in summer.

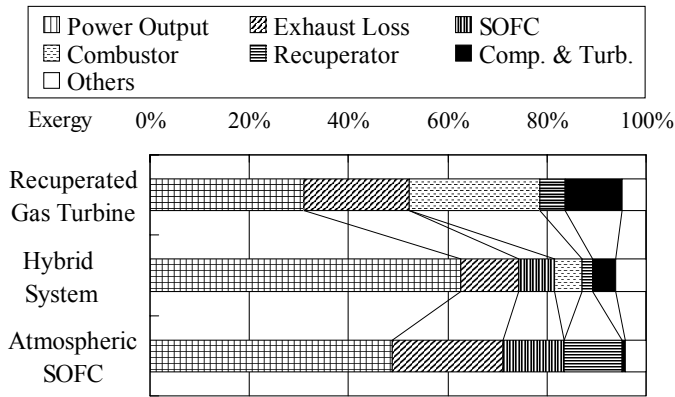


Fig. 2 Breakdown of exergy consumption

Table. 2 Effect of power output capacity

	Recuperated GT		Hybrid System	
	30	300	30	300
Total Output (kW)	30	300	30	300
Pressure Ratio (-)	5.0	6.0	5.0	5.0
Compressor Adiabatic Efficiency (%)	75	79	73	76
Turbine Adiabatic Efficiency (%)	85	87	80	86
GT Mechanical Efficiency (%)	97	98	95	97
Turbine Inlet Temperature (°C)	1250			
Fuel Consumption per 1kW Output (kW)	2.65	2.42	1.54	1.47

Table. 3 Effect of intake air temperature

	Recuperated GT		Hybrid System	
	15	30	15	30
Total Output (kW)	30			
Compressor Inlet Temperature (°C)	15	30	15	30
Pressure Ratio (-)	5.0	5.0	5.0	5.0
Compressor Adiabatic Efficiency (%)	75	73	73	73
Turbine Adiabatic Efficiency (%)	85	80	80	80
GT Mechanical Efficiency (%)	97	95	95	95
Turbine Inlet Temperature (°C)	1250			
Fuel Consumption per 1kW Output (kW)	2.65	2.77	1.54	1.57

## 5. STRATEGY FOR HYBRID SYSTEM DESIGN

Some of the design parameters of a hybrid system are flexible, while the others cannot be changed widely. For example, once the output power is set, the compressor and turbine efficiencies are given, whereas we can choose the pressure ratio rather freely. Here, the effects of cell temperature and of TIT on the cycle performance are shown. Moreover, general strategy to set cycle parameters, such as recuperator temperature effectiveness, S/C ratio, pressure ratio and TIT, is also shown below.

### 5.1 Effects of Cell Temperature and TIT

While we study the effects of cell temperature and TIT, the other variables are fixed. The generation efficiency is shown in Fig. 3. First, it is clear that higher cell temperature gives higher

efficiency. However, unlike a conventional gas turbine, the hybrid system's efficiency is decreased with increasing TIT at a fixed cell temperature. We should remember that higher cell temperature implies higher cost and requires longer time to start up. Therefore, we fix the cell temperature at 1000°C, that is the limit for the state-of-the-art SOFC technology, in the following discussions.

Figure 4 shows the total heat input (thicker line) and the SOFC heat input (thinner line). As a result, the heat input to the

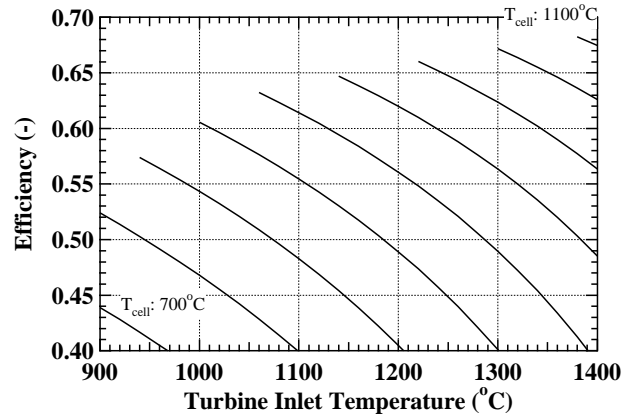


Fig. 3 Power generation efficiency (LHV)

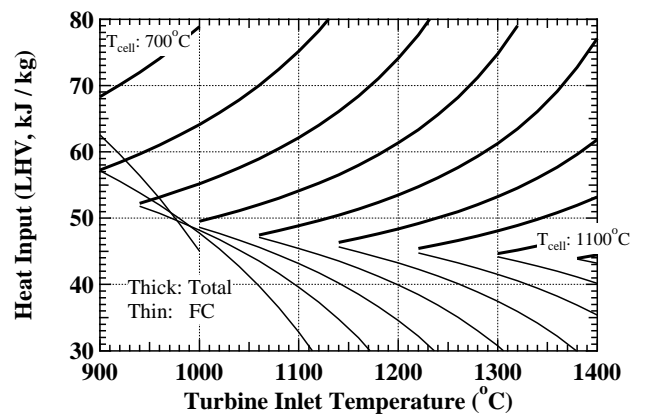


Fig. 4 Heat input

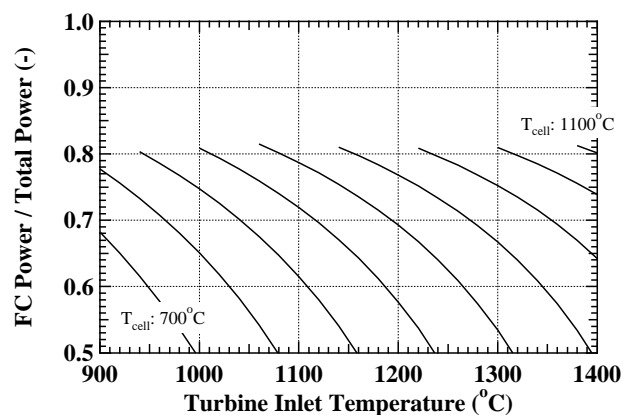


Fig. 5 Fraction of SOFC output

combustor is given as a difference between them. Large direct heat input to the combustor causes high TIT. Thus, it is found that large direct heat input to the combustor reduces total generation efficiency, because the heat input to SOFC is utilized more effectively than is the combustor. This is supported by Fig. 3, in which high TIT or large heat input to the combustor gives low generation efficiency.

It is also evident that there is a certain TIT for each cell temperature with zero heat input to the combustor. We can not lower TIT beyond the temperature that is equivalent to zero heat input.

The ratio of SOFC output to total output is shown in Fig. 5. Higher cell temperature gives higher SOFC fraction, while higher TIT gives lower SOFC fraction. At the lowest TIT for each cell temperature, SOFC fraction is approximately 80%.

### 5.2 Recuperator Effectiveness

We estimate the effect of recuperator temperature effectiveness on the cycle performance. High recuperator effectiveness leads to high cathode inlet temperature. Thus, the fuel input to SOFC for a unit amount of air is reduced if the cell temperature is fixed. This is confirmed by the decrease in oxygen utilization in SOFC (Fig. 6). Small fuel input to SOFC causes residual gas such as methane, hydrogen and carbon oxide decrease (Fig. 7). Less

residual gases lower TIT. Therefore, higher recuperator effectiveness causes lower TIT for a fixed heat input to the combustor (Fig. 8). The generation efficiency (Fig. 9) is increased if the recuperator effectiveness is increased, when TIT is decreased or the fuel input to the combustor is fixed.

### 5.3 Steam Carbon (S/C) Ratio

S/C ratio is defined as the ratio of the mole fractions of steam and methane at the reformer inlet. The S/C ratio determines the amount of anode recirculation.

When we increase the S/C ratio with other parameters fixed, the amount of anode recirculation or that of anode inlet gas is increased. This results in the reduction of the fuel input to the SOFC at a fixed SOFC outlet temperature. In the same manner, as the recuperator effectiveness is increased, a high S/C ratio causes low TIT where the fuel input to the combustor is fixed (Fig. 10). Figure 11 shows the generation efficiency. Increase in S/C ratio reduces the generation efficiency with fixed TIT. But, if S/C ratio decreases with fixed fuel input to the combustor, reduction of the generation efficiency is not remarkable.

We can decrease the turbine exit temperature with higher S/C ratio and lower TIT. If S/C ratio is increased, for example, from 3.0 to 4.0, minimal TIT decreases from 1210°C to 1165°C. Simul-

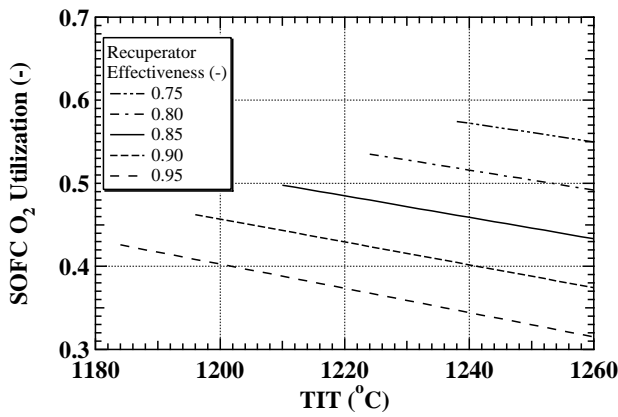


Fig. 6 O<sub>2</sub> utilization in SOFC

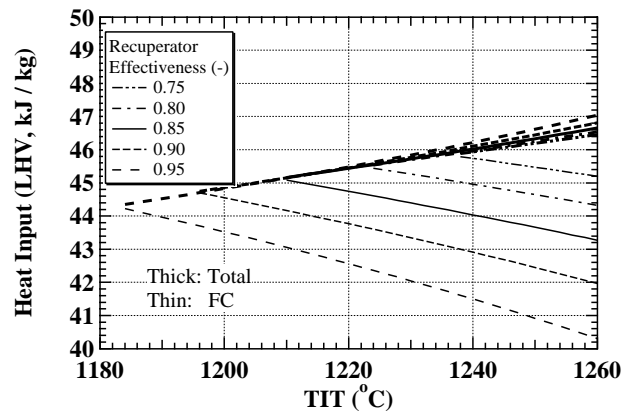


Fig. 8 Heat input

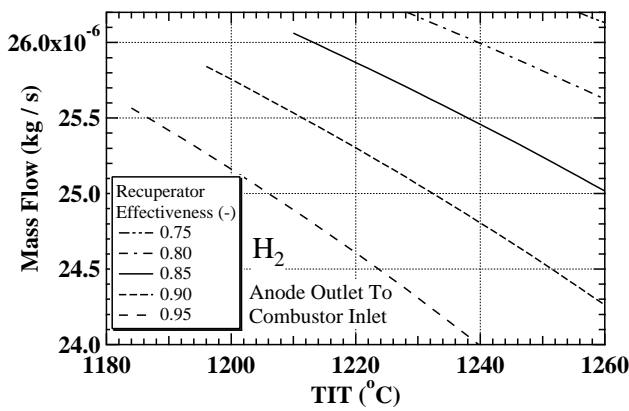


Fig. 7 H<sub>2</sub> mass flow into combustor

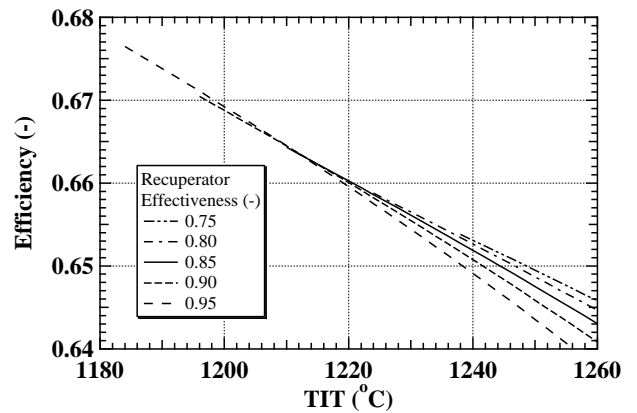


Fig. 9 Power generation efficiency (LHV)

taneously, the turbine exit temperature is lowered from 845°C to 810°C. There is a maximum temperature for the use of metallic recuperators between 850°C and 900°C. Beyond this temperature, we should utilize ceramic recuperators with much more cost<sup>(2)</sup>. High S/C ratio may enable us to adopt metallic recuperators, and this is favorable even if some reduction of the generation efficiency is considered.

### 5.4 Pressure Ratio

So far, it is clarified that high recuperator effectiveness is preferable as well as a high S/C ratio moderates the requirement of the temperature increase. We set recuperator effectiveness at 95%<sup>(2)</sup>, and S/C ratio at 4.0 in the following discussions.

To optimize the system pressure and TIT, we performed cycle calculation. Figure 13 shows the effect of pressure ratio on generation efficiency. The maximum efficiency is approximately 67% (LHV) at pressure ratio 4.5 with zero direct heat input to the combustor. However, variation of the maximum generation efficiency from pressure ratio 2.5 to 6.0 is not remarkable.

Higher pressure ratio reduces the turbine outlet temperature and cathode inlet temperature. Low cathode inlet temperature causes high SOFC fuel input. Contrary to the effects of recuperator effectiveness and S/C ratio, higher pressure ratio gives higher TIT. Thus, it is clear that TIT depends on the pressure ratio. We plot heat input in Fig. 14 to determine TIT. Smaller direct fuel input to the combustor is preferable, but a certain amount of fuel input is necessary to burn anode residual gases completely. Assuming that the heat input to the combustor is 1.5kW, we can estimate TIT = 1150°C in Fig. 14 for a pressure ratio of 4.0. At this point, the ideal cell voltage is 0.863V, and the net cell voltage is 0.704V. This difference of 0.159V is due to polarization (Fig. 15).

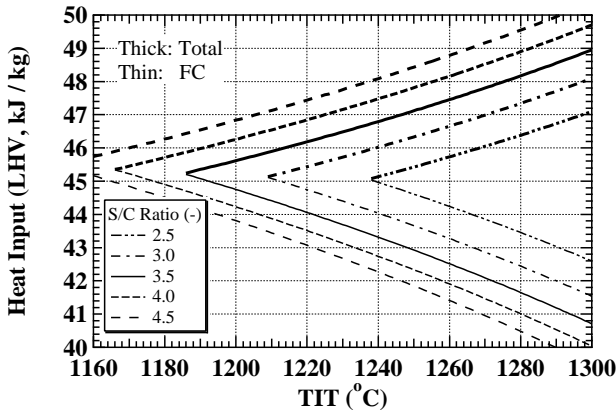


Fig. 10 Heat input

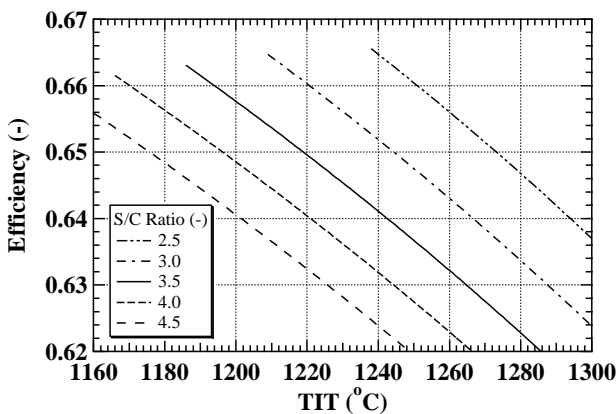


Fig. 11 Power generation efficiency (LHV)

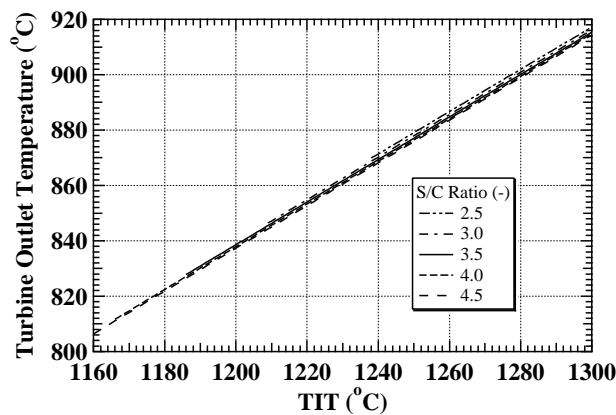


Fig. 12 Turbine outlet temperature

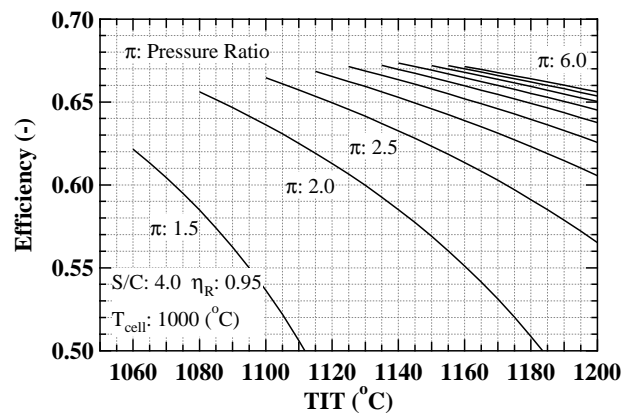


Fig. 13 Power generation efficiency (LHV)

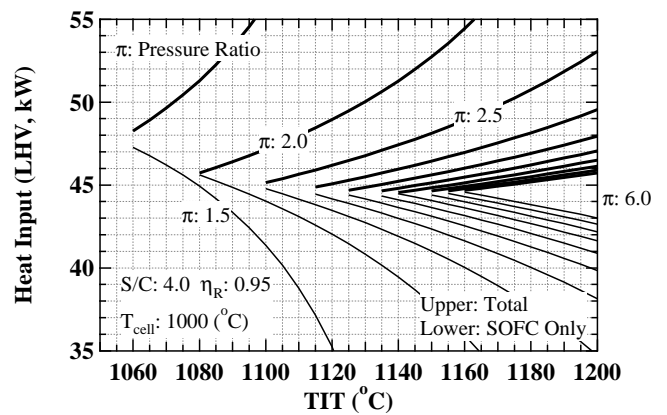


Fig. 14 Heat input

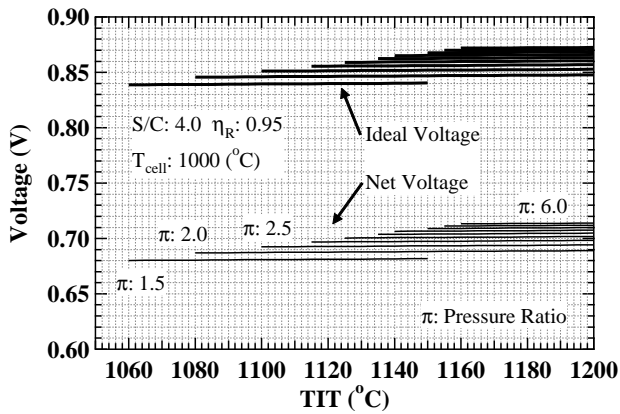


Fig. 15 Cell Voltage

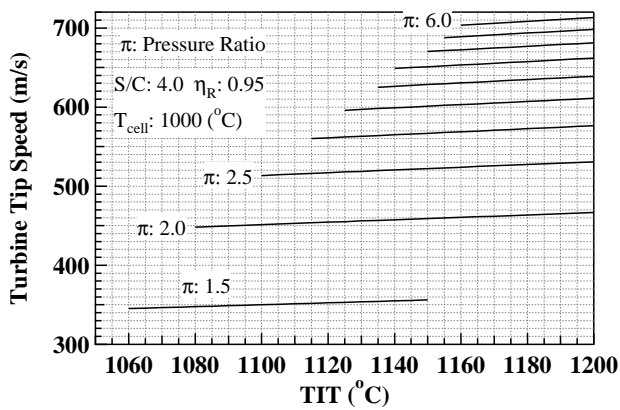


Fig. 16 Estimated turbine tip speed

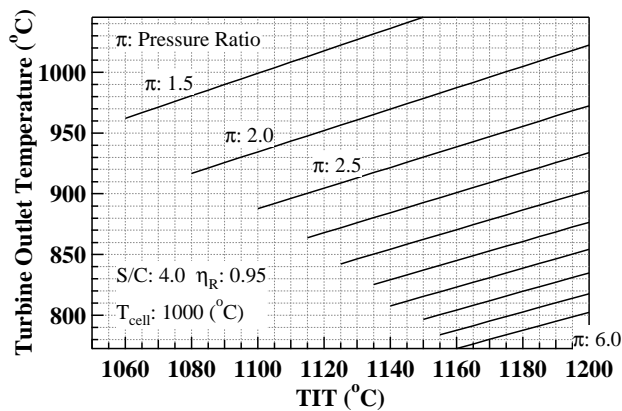


Fig. 17 Turbine outlet temperature

We should also account for the aspects of strength and high-temperature durability of materials. Lower turbine tip speed, which causes lower turbine stress, is desirable. Low turbine outlet temperature is also preferred because it enables us to utilize inexpensive materials for recuperators. The turbine tip speed (Fig. 16) increases as the pressure ratio increases contrary to turbine outlet temperature (Fig. 17). Therefore, we should compromise them. For pressure ratio 4.0 and TIT 1150°C, the turbine tip speed is approximately 630m/s and the turbine outlet temperature is 840°C. These values should be technologically feasible in the foreseeable future.

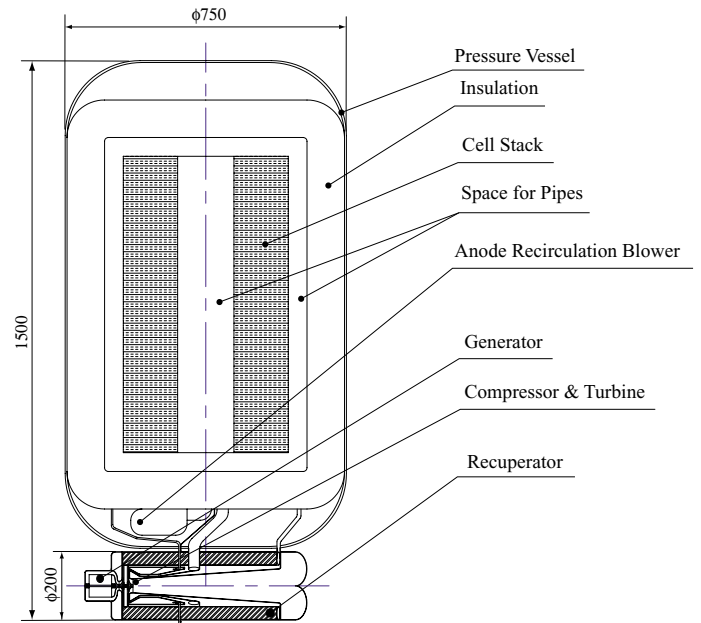


Fig. 18 Conceptual design

## 6. CONCEPTUAL DESIGN

We performed a conceptual design of a 30kW μGT-SOFC hybrid system for the best possible case. As mentioned above, we set pressure ratio at 4.0, TIT at 1150°C, S/C ratio at 4.0, and recuperator effectiveness as 0.95. The total area of the cell is estimated from the cell current required divided by the current density. The thickness of the cell is estimated from an existing research model of SOFC<sup>(9)</sup>, whose thickness is 5mm. We consider 3mm for reformer and supports. Then, the thickness of a single cell is 8mm. The conceptual design is shown in Fig. 18. The overall size is about of a refrigerator for home use.



## 7. CONCLUSIONS

Cycle calculations of  $\mu$ GT-SOFC hybrid systems and a conceptual design of a hybrid system for the best possible case are carried out. The following conclusions are derived:

- (1)  $\mu$ GT-SOFC hybrid system has potential to achieve high efficiency as high as 65% (LHV), even for a small system of 30kW.
- (2) Hybrid systems have higher generation efficiency than that of traditional recuperated gas turbines or atmospheric SOFC, because the exergy loss of combustion and exhaust is much lower.
- (3) Output capacity and ambient temperature have smaller influence on the hybrid system than on traditional recuperated gas turbine systems.
- (4) Lower amount of direct fuel input to the combustor increases generation efficiency.
- (5) Amount of direct fuel input to the combustor depends not only on TIT, but also on other parameters such as recuperator effectiveness, S/C ratio and pressure ratio.
- (6) Higher recuperator efficiency gives higher generation efficiency and lower TIT.
- (7) Higher S/C ratio lowers TIT and turbine outlet temperature, although it causes a little decrease in generation efficiency.
- (8) A certain pressure ratio gives the maximum generation efficiency, but variation of generation efficiency near the optimal pressure ratio is not remarkable.
- (9) The diameter of 30kW  $\mu$ GT-SOFC hybrid system is estimated as about 750mm, and its height 1500mm.

## ACKNOWLEDGMENTS

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