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TIMBER UNDER REAL FIRE CONDITIONS – THE INFLUENCE OF OXYGEN CONTENT AND GAS VELOCITY ON THE CHARRING BEHAVIOR

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ABSTRACT: As for any building material, verification of fire resistance is mandatory for timber structures. While non-standard fire design for steel members has long tradition, the corresponding possibilities for timber members are limited. Reasons for this can be found in the degree of complexity of the material and the limited research done in the field. This paper summarizes outcomes of tests, investigating the influences on the charring behaviour of timber by varying the oxygen content and the gas velocity in the compartment. Results show that charring is depending on the fire compartment temperature, but, results show further that at higher oxygen flow, char contraction was observed affecting the protective function of the char layer. In particular, in the decay phase char contraction should be considered which may have a significant impact to Performance Based Design using non-standard temperature fire curves where the complete fire history including the cooling phase has to be taken into account.

KEYWORDS: Timber, Char contraction, Charring, Non-standard fire, Fire resistance

1 INTRODUCTION AND BACKGROUND

Fire design of structural timber members is done in Europe according to rules given in the fire part of Eurocode 5, EN 1995-1-2 [1]. Design models specified herein are usually developed for standard fire exposure [2]. Very limited models are available for non-standard fires. In general, design of timber members considers (i) the reduction of the cross-section (CS) by char as well as (ii) the reduction of strength and stiffness of the residual cross-section (RCS). The Reduced Cross-Section Method, or Effective Cross-Section Method addresses (i) by a time depending charring depth and (ii) by a so called zero strength layer. The latter is not content of this study. While the charring rate in standard fires according to ISO 834 or EN 1363-1 [2] is reported to be constant [3], charring rates for non-standard fires varies and existing rules may be questioned.

The origin of this study is based on observations in fire resistance tests following the standard fire exposure

(defined time-temperature and pressure curves as well as minimum oxygen content of 4 % [2]) in different furnaces where different charring rates were observed for the same product. An additional motivation for this study is observations of fire accidents where different RCS of solid timber members inside a fire compartment depending on the air flow [4].

2 TEST PROGRAM

In total 14 tests with various test conditions were performed in a custom made, gas fired furnace. The tests lasted 60 minutes after ignition. After each test the timber specimen was removed from the furnace and any burning was extinguished with water; this procedure took less than 30 seconds to allow for accurate evaluation of the RCS and the char layer depth.

2.1 MATERIAL

Tests were performed with CLT (Cross Laminated Timber) beams (150 mm x 150 mm x 1730 mm, C24

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European wood grade) with $12 \% \pm 1 \%$ equilibrium moisture content. The five layer product (top layer 42 mm) is glued with PUR (Polyurethane) adhesive and all specimens were stored in a conditioning room before testing. CLT was used as it was aimed for a homogeneous material with limited number of defects. The timber members with a width of 150 mm were insulated on both sides with stone wool (initial thickness 45 mm and density of 35 kg/m^3) to evoke one dimensional heat exposure and charring of the surface exposed to fire. In four length positions (CS A to D) the progression of char, temperatures and the oxygen content were recorded during the tests (measurement stations). Temperature measurements were done by means of internal thermocouples type K with welded junction (wire $\varnothing 3 \text{ mm}$) inserted in horizontal bore holes $\varnothing 1.75 \text{ mm}$; wires were inserted so that the first 50 mm after the hot junction were parallel to the Isotherms.

2.2 TEST EQUIPMENT

The equipment used in the tests is shown systematically in Figure 1. The test set-up **Figure 2** consisted of (i) a gas supply unit in which a defined gas volume and the oxygen content of the supplied gas were controlled by means of mixing ambient air with nitrogen or oxygen, (ii) a heating device where the supply air was mixed with burning Gasol (95 % Propane and 5 % Butane), (iii) a fire compartment (insulated steel channel 200 mm x 160 mm x 2000 mm) and, (iv) an exhaust unit (part of the fire lab). Three sides of the fire compartment were designed to function as a large plate thermometer (PT) [2] comprising a steel plate at the exposed side and ceramic fibre insulation (double layer in order to address the temperature differences) at the unexposed side. Temperatures were measured at sections A to D on the top and side of the fire compartment. Tests were performed in low under-pressure (ca. 10 Pa) to provide a good working environment in the laboratory. The air velocity was controlled in the inlet (cold condition) as well as in the fire

compartment using different measurement techniques, an anemometer as well as a Pitot tube coupled with a temperature measurement [5]. Both measurements were in a reasonable good agreement. The oxygen content of the air was measured with a paramagnetic and an electrochemical method in five CSs, at the beginning of the fire compartment (CS zero) and the sections A to D distributed over the length of the specimen. The sample gas was taken from the compartment by placing the probe head (electrochemical method) and the sample pipe (paramagnetic method) respectively in contact to the timber surface. By means of openings at the side of the probes, the sample gas was taken from distances of about 3 to 10 mm from the surface.

Data measurements (pressure, temperature, gas velocity, oxygen concentration) were recorded with intervals of 1 to 60 seconds. The gas flow (ambient air, nitrogen, oxygen, Propane) was adjusted manually and the Propane gas flow (kg/h) was recorded manually when the set point was changed.

2.3 TEST CONDITION

All fire tests aimed for a quick initial fire compartment temperature rise and a constant temperature during the test. During the early testing phases, different temperatures of the compartment sections A to D were measured. Additionally to the PT temperature, the gas temperature inside the fire compartment was measured using small thermocouples. The gas temperature was further used to determine the gas velocity in the fire condition in combination with a Pitot tube [5]. The temperatures in the compartment followed an exponential temperature increase, in later phases of each test a temperature plateau between about 700°C and 900°C was observed.

Gas velocities were varied between about 1 and 15 m/s in the fire compartment and the oxygen between 5 and 15 % (percentage by volume). This was accomplished by setting the temperature level by means of the Propane

Figure 1: Schematic CS view of the test set-up

burner at ambient air flow and then adjusting the mixture of ambient air as well as the gases nitrogen and oxygen from the gas tanks in order to reach the set point for the oxygen level together with the gas velocity, see Table 1. This procedure of firing the fire compartment was a looped process.

Table 1: Test program (excluding calibration tests)

Test	6	7	8	9	10	11	12	13	14
Oxygen [%]	5	15	5	15	15	15	10	15	15
v [m/s]	1	10	1	3	3	3	10	10	10
Comparison	(a)	(b)	(a)	(b)		(b)	(c)	(b)	(b)
		(c)						(c)	(c)

3 RESULTS

The documentation comprises the following recordings: (a) gas concentrations, (b) pressure, (c) gas velocity, (d) temperature and (e) documentation of the specimen (RCS, char layer depth) after the tests. For the evaluation of charring depth in longitudinal direction, residual longitudinal-sections at the centre lines of the timber specimens were documented, see Figure 2. **Error! Reference source not found..**

Figure 2: RCS (length profile), and total depth after the fire Test 8 (1m/s gas velocity, 5 % O₂)

To evaluate the RCS at the corresponding temperature measurement stations (CS A to D), as well as effects of the limited CS width of the beams, images of the RCS were taken at the measurement stations, see Figure 3. To follow the progress of charring within the CS during the test the movement of the 300°C isotherm, assumed to be equal to the char line [1], was evaluated using the internal thermocouples for all specimens and all sections. The results of the char progression in the corresponding depths of the thermocouples (6 to 42 mm; step +6) are presented for Tests 06 and Test 08 (gas velocity of 1 m/s and 5 % oxygen content) in Figure 4.

Figure 3: Remaining CS (including the char layer) and residual CS in the length allocated to CS B thermal exposure (measurements at 700 mm)

Figure 4: Estimated charring depth by means of the internal thermocouples of Test 6 and 8 in CS C and D.

3.1 Comparison of test results

Comparison was done by evaluating test results of similar tests varying one parameter, i.e. (a) the temperature (b) the oxygen content and (c) the gas velocity, see Sections 3.1.1 to 3.1.3.

3.1.1 Comparison (a) – Variation of the temperature

In Tests 06 and 08, see Table 1, the gas velocity as well as the oxygen content were set to the same value, 1 m/s and 5 % O₂. The temperature plateau was deviating as the burner effect was increased in test 08. In Figure 4, it is shown that the test done with a higher temperature shows slightly higher charring rates. The difference of the charring rate is 0.12 mm/min in CS C and 0.09 mm/min in CS D where slightly lower compartment temperatures (PT temperatures) were measured. As a consequence, the RCS was less for higher temperature, although the total CS depth was observed to be about the same, i.e. 150 mm, which means that no char contraction occurred, see Figure 5.

Figure 5: RCS (length profile) and total depth after Tests 06 and Test 08 (1m/s gas velocity, 5% oxygen).

3.1.2 Comparison (b) – Variation of the gas velocity

The determined charring depth of Test 07, 13 and 14 ($v=10$ m/s) were compared to Test 09 ($v=3$ m/s). All tests within this comparison (b) had an oxygen of 15 %. Besides the oxygen content, the temperature was the same for all of the four tests in the steady phase of the temperature-time curve (plateau), compare specifications in Table 1. A difference of 12 mm thickness in the char layer was observed in CS A comparing the high gas velocity tests and the low velocity test, see Figure 6. Thus, the maximum difference of the charring rate (in CS A) can be assessed to be about 0.29 mm/min, see Figure 6. It was concluded that timber members exposed to (i) high gas velocity and (ii) high oxygen content, i.e. substantial mass flow of oxygen lead to higher charring rates.

Figure 6: Maximum deviation of the charring depth estimated by means of the internal thermocouples of Test 07, 13, 14 against Test 09 in CS A.

Further, analysis of the RCS showed that the high gas velocity combined with the high oxygen content lead to a reduction of the char layer thickness. This result was more distinct in CS A than at the end of the length profile of the specimen.

3.1.3 Comparison (c) – Variation of the oxygen content

In the last comparison, Tests 07, 13 and 14 (O_2 content 15 %) were compared with Test 12 (O_2 content 10 %) to study the dependence of the charring rate on the oxygen content. As in the previous comparisons (a) and (b), the other two parameters (in this case gas velocity and temperature) were set to the same value in these tests, see Table 1.

Figure 7: Maximum deviation of the charring depth estimated by means of the internal thermocouples of Test 07, 13, 14 against Test 12 in CS A.

In CS A, as shown in Figure 7, the char layer depth of the high oxygen content tests is 6 mm larger than for the low oxygen content test.

Figure 8: Total depth (length profile) of specimens of Tests 07, 13, 14 against Test 12 in CS A

Thus, also corresponding charring rates are different. This may be traced back to the reduction of the char layer thickness as well as the glowing combustion. Tests with a high oxygen content showed higher charring rates than Test 12. The maximum difference in CS A is 0.16 mm/min.

The RCS and the total CS are lower for the tests with a high oxygen percentage (Test 07, 13 and 14) if compared with the low oxygen content test over the entire length of the beam, see also Figure 8.

4 ANALYSIS OF THE EFFECT OF GAS VELOCITY

To analyse the effect of gas velocity and/or the mass flow of oxygen, thermal simulations were performed. The aim of the simulations was to estimate the effect of the gas velocity in comparison to temperature on the specimen. In the following, two tests (Test 09 and Test 14) are compared where equal temperature time curves were achieved. The oxygen content was set to 15 % and the gas velocity was 3 and 15 m/s respectively. For the simulations it is assumed that the effective material properties for standard fire are valid which are given in [1]. This assumption is justifiable due to the limited deviations of the temperature exposure from the standard fire exposure.

Figure 9: Length profiles of RCS and total depth of specimens of Test 9 and Test 14 in CS A.

As for the other tests, the temperature exposure showed almost a logarithmic temperature rise (ca 30 min) up to a plateau at about 800°C; the tests lasted 60 min. Fire exposures (mean PT temperatures) are specified in Figure 10. Comparison of the specimens after the test showed different CS depths for tests with respect to the total remaining CS (total depth including the char layer) and the RCS (virgin or uncharred wood). The length profile was observed to be most steady in CS B at about 700 mm length position which is further evaluated. It is assumed that the compartment temperature measurements (PT measurements) of any CS are valid for a length section of about 500 mm. In the following, the RCS and temperature measurements in CS B are further analysed, the corresponding observed profiles of the RCS and the total cross-section depth is given in Figure 9.

Figure 10: Temperature in tests no. 09 and 14 and the standard fire exposure

4.1 SIMULATION OF CHAR CONTRACTION

One dimensional finite element temperature simulations were performed with a section of 1 mm width. Thermal simulations were performed using the software SAFIR [6]. Standard values for surface emissivity, $\varepsilon = 0.8$ and the convective heat transfer coefficients, $\alpha_{unexp} = 9$ and $\alpha_{exp} = 25 \text{ Wm}^{-2}\text{K}^{-1}$, were used. The sides of the 1-D section were assumed adiabatic and the unexposed side held at 20°C. Simulations used the effective material properties for density, heat capacity and heat conductivity available in EN 1995-1-2 [1] although this values are intended for standard fire exposure. This approach was found to be reasonable as the deviation of the actual fire exposure from the standard fire temperature time curve is limited, see Figure 11.

Figure 11: Simulated surface temperatures and temperature at a depth of 40 mm for different fire exposures with (case ii) and (iii)) and without (case i)) char contraction.

At standard fire exposure, charring depth after one hour can be expected to be about 40 mm according to the simplified, linear charring rules in [1]; this value is supported by numerous studies. In the following, this depth was used as a reference depth at 60 min, i.e. the temperature rise in 40 mm depth is analysed with respect to different assumptions to investigate char contraction. In Test 14, char contraction was observed to be about 10 mm after 60 min.

Char contraction was assumed to occur linearly which was realized in the simulations by deleting elements in equal steps of 6 min. Thus, for simulations, every 6 min the calculation was stopped, one element (1 mm x 1 mm) deleted and simulations were re-started using the before obtained temperature field for the remaining elements. The following cases were investigated:

- (i) Standard fire exposure and no char contraction as basis for the following comparison.
- (ii) Standard fire exposure and char contraction to document the importance of the char layer as insulating layer.
- (iii) Actual fire exposure and char contraction.

4.2 SIMULATION RESULTS AND CONCLUSION

Results for case (i) show lead charring of 40 mm at about 63 min which is only slightly later than the conservative rules of EN 1995-1-2 [1], see Figure 11. In corresponding fire tests performed in this study, the resultant charring depth was 35 to 38 mm after 60 min.

Results for case (ii) show that the temperature increase in 40 mm deviates significantly after 30 min resulting in charring at this depth about 8 min earlier than in case (i). This indicates that the char layer provides significant protection of the virgin wood in the inner part of the CS. The significance of the char layer has been investigated in many tests with cross-laminated timber where char ablation (e.g. for Cross-Laminated Timber) lead to a significantly increased charring rate.

Results for case (iii) show that considering the slightly lower fire exposure measured in Test 14 charring would not reach 40 mm depth. This is in contradiction to the observations where the RCS was significantly below 110 mm (corresponding 40 mm charring depth) which would be more appropriate for standard fire exposure.

Considering the actual difference of the char layer depth (10 mm) and of the charring depth (42 and 35 mm) it can be assumed that the only varying parameter, the gas velocity, is finally responsible for the char oxidation and subsequently the smaller RCS. It is assumed that the actual gas velocity in combination with the oxygen content leads to glowing combustion which causes char contraction. It may be concluded that the effect is corresponding to an increased fire compartment temperature of up to 200°C.

5 ESTIMATION OF THE HEAT RELEASE BY THE TEST SPECIMEN

When estimating the fire development in a compartment, the fire load of timber structures has moved into the focus of many authorities and researchers especially since solid timber products, e.g. CLT, appeared on the market. With

the actual test set-up, it was possible to estimate the contribution of the test specimen by two alternative methods. Firstly, (A) based on the charring rate (charring depth), and secondly, (B) by means of the oxygen consumption.

5.1 ESTIMATION OF THE HEAT RELEASE BY MEANS OF CHARRING (Method A)

Cone calorimeter tests have previously been conducted to determine the heat release rate of different materials under certain exposures of constant incident heat flux. In this test, a specimen is exposed to a homogeneous incident heat flux. It can be concluded that the following ratio is valid:

$$HRR : \beta = HR : d_{char} \quad (1)$$

where

HRR is the heat release rate,

HR is the heat release,

β is the charring rate and

d_{char} is the charring depth.

As a basis for a compartment fire model, a relationship between the charring rate and the heat release rate was determined from cone calorimeter test results. Investigations of heat release rates and corresponding charring were investigated in [7].

Figure 12: Heat release rate of untreated timber under 75kW/m² exposure; reproduced from [7]

At 75 kW/m² incident radiant heat flux, a heat release of 5400 kJ/m² per millimetre of charring depth, for charring depths exceeding 10 mm was determined. Results are specified in Section 5.3.

5.2 ESTIMATION OF THE HEAT RELEASE BY MEANS OF THE OXYGEN CONSUMPTION (Method B)

The estimation of the heat release rate can be performed from the oxygen consumption using calorimetry [8]. It can be assumed that the heat release per consumed mass of oxygen is 13.1 MJ/kg for most bio-based materials.

In the test setup, the oxygen concentration was measured in CS zero, A, B, C and D. Further, the gas velocity in the

compartment was determined and was assumed to be constant. The mass of oxygen passing each section can be determined in a similar way as is done using a cone calorimeter. Results are specified in Section 5.3.

5.3 RESULTS OF THE HEAT RELEASE ESTIMATION

The calculated heat release rates according to the methods presented in Sections 5.1 and 5.2 are in rough agreement, see Figure 13. However, the results are not accurate enough to explain differences in heat release rates between tests, as indicated in Table 2.

Table 2: Estimated heat release rate of Test 14 (15m/s, 15% O₂) and Test 09 (3m/s, 15% O₂)

	Heat release rate (kW/m ²)		
	Method 1		Method 2
	Section B	Section C	Between section B and C
Test 14	55.9	48.7	48.5
Test 9	44.5	52.7	61.7

Figure 13: Estimated heat release rates using method A and B

6 METHODS FOR SAMPLE GAS AND GAS VELOCITY ANALYSIS

The study used redundant systems to determine the (i) gas velocity and the (ii) oxygen content.

The gas velocity was estimated using a Pitot tube [5] combined with a small thermocouple wire. Further, the gas velocity was determined using an anemometer at the gas inlet (after mixture of ambient air, oxygen and nitrogen) adding to the effect of the combustion gases of the Propane burners. The estimation was performed considering the thermal expansion of the gases. The latter was done using the gas temperature of the fire compartment. The combustion of Propane increased the inaccuracy as a combustion factor had to be introduced in order to take into account for the incomplete combustion. Differences between the calculation (anemometer) and more direct measurements (Pitot tube) were most visible

at high gas velocities and can be specified to about 20%. It was concluded that Pitot tube measurements are more reliable for measuring the gas velocity in fire compartments.

Compartment gas analysis was performed using a hand held electrochemical gas analyser and a paramagnetic method. The electrochemical gas analyser offered in-situ analysis of several parameters (gas types) which gave similar results ($\pm 1\%$) than the more expensive paramagnetic analysis which had to extract sample gas from the compartment. In contrast to cone-calorimetry, it seems that the electrochemical electrodes provide sufficiently accurate results for the test set-up presented here [9].

7 DISCUSSION

7.1 Char contraction

Earlier studies reported char contraction but the observed phenomenon could not be explained [10,11]. In contrast to these two studies, in this study char contraction was investigated in a closed compartment with a controlled gas flow. In this study, the boundary conditions in the combustion compartment were well defined. Results show that a variation of the fire compartment environment may lead to a char contraction of about 10 mm after 60 min which is in line to very recent studies [12].

Comparison (c), presented in Section 3.1.3, showed that a fire compartment oxygen content of 15 % lead to char contraction while using oxygen content of 10 and 5 % gave no significant effect. It seems like there is a limit for significant glowing combustion between 10 and 15 %. This is in line with another study of calorimetric experiments where the critical oxygen limit was found to be 14 % [13].

For an oxygen content 15%, the gas velocity influenced the char contraction, about 10 mm in difference, see Figure 9. This can be explained by the increased mass flow (turbulence) and thereby the enlarged transport of oxygen to the combustible specimen's surface. Thus, the temperatures in the virgin wood were different depending on the gas velocity as shown in Section 3.1.2. This is obvious by following the progression of the 300°C isotherm, assumed to be the char line [1], as shown in Figure 6.

Based on Comparisons (b) and (c), presented above, it seems that char contraction is rather a result of the gas velocity of a distinct concentration above a critical limit, than the oxygen content only. This explains, why no clear results were found in other studies where the boundary conditions were not sufficiently defined [10,11]. To assess the effect on timber member temperature of the char contraction (in combination with glowing combustion) simulations were performed, see Section 4. Simulations with standard fire exposure and char contraction (case (ii)) show that the char layer works as effective protection of the virgin wood. It was further shown (case (iii)) that the temperatures at the char line (target temperature 300°C at a depth of 40 mm as observed from the test results) would still be very low (ca. 180°C) but glowing combustion induced by gas velocity accounts for the significant difference, see Figure 11.

Two methods were used to estimate the heat release rate caused by the combustible test specimen in different testing environments. For cone calorimeter tests both test methods were compared in an earlier study [9]. The methods give results between ca. 45 and 60 kW/m², however, heat release rates over the time show considerable inconsistencies, see Figure 13. Further, an increased heat release rate for Test 14 where significant glowing combustion was determined (compared to Test 09, see Section 4) could not be confirmed, see Table 2.

8 CONCLUSIONS

In the literature, the char layer development is considered to be a crucial part of analysis of timber members exposed to fire as it provides protection of the virgin wood. In more recent studies concerns are raised that the charring adds to the fire load when glowing. In this study, the focus was (a) the contribution of the char glowing combustion to the overall heat release and (b) the glowing effect on the residual cross section. In order to study these effects, the fire compartment environments were simulated by varying (i) the compartment temperature, (ii) the gas velocity and (iii) the oxygen content. In fires gas velocities up to 15m/s are reasonable while during a fully developed fire the oxygen content of the compartment gas is very low for ventilation controlled fires. However, as performance based design requires the consideration of an entire fire, the decay phase should also be considered.

In this study, it was proven that the temperature has a significant influence on the charring behaviour of a timber member. Moreover, it was shown that glowing combustion influences the char layer (char contraction) for oxygen contents above about 15 %. The effect becomes more significant when the gas velocity is increased.

Following this conclusion many fire resistance tests simulating entire including the cooling phase, may be questioned as the ventilation conditions (oxygen content, gas velocity) in the cooling phase were not controlled and not documented. Studies that have investigated different ventilation conditions are very rare but available literature indicate significant differences [14].

In performance based design, the char contraction may be taken into account in the decay phase depending on the ventilation conditions due to temperature differences, e.g. in stair cases.

Char contraction was estimated only after the test in lack of methodologies to measure this value during the test. Thus, it cannot be stated in which phase char contraction occurs. In future tests, the char contraction should be investigated further.

Considering the results presented for the heat release, no dependency on the glowing combustion of the timber member on the oxygen mass flow could be determined, see Table 2.

In future studies the methods presented in this paper should be improved the estimations of the heat release rate. Further, the time of heat exposure should be extended and the decay phase further evaluated.

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