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# Computer-aided working-fluid design, thermodynamic optimisation and technoeconomic assessment of ORC systems for waste-heat recovery

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## Abstract

The wider adoption of organic Rankine cycle (ORC) technology can be facilitated by improved thermodynamic performance and reduced costs. In this context the power system should be evaluated based on a thermoeconomic assessment with the aim of improving economic viability. This paper couples the computer-aided molecular design (CAMD) of the working-fluid with thermodynamic modelling and optimisation, in addition to heat-exchanger sizing models, component cost correlations, and a thermoeconomic assessment. The proposed CAMD-ORC framework, based on the SAFT- $\gamma$  Mie equation of state, allows the thermodynamic optimisation of the cycle and working-fluid in a single stage, thus removing subjective and pre-emptive screening criteria that would otherwise exist in conventional studies. Following validation, the framework is used to identify optimal working-fluids for three different heat sources (150, 250 and 350 °C), corresponding to small- to medium-scale applications. In each case, the optimal combination of working-fluid and ORC system is identified, and investment costs are evaluated. It is observed that fluids with low specific-investment costs (SIC) are different to those that maximise power output. The fluids with the lowest SIC are isoheptane, 2-pentene and 2-heptene, with SICs of 5,620, 2,760 and 2,070 /kW respectively, and corresponding power outputs of 32.9, 136.6 and 213.9 kW.

*Keywords:* organic Rankine cycle; ORC; computer-aided molecular-design; CAMD; group contribution; SAFT- $\gamma$  Mie; technoeconomic.

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## 1. Introduction

Despite growing interest in improving energy-efficiency to reduce fossil fuel consumption and our impact on the environment, there remains a significant amount of waste heat that is currently rejected to the atmosphere. Of the technologies that can be considered for waste-heat recovery, the organic Rankine cycle

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5 (ORC) is one of the most promising candidates, and is suitable for converting low- and medium-grade waste  
6 heat, typically at temperatures between 80 and 400 °C, into electricity [1, 2].

7 One of the most important components of an ORC is the working fluid, and the fluid selected can affect  
8 performance, component design, size, cost and operational procedures. However, with increasing concerns  
9 over global warming and air pollution, certain fluids such as chlorofluorocarbons have already been phased  
10 out, whilst fluids such as hydrochlorofluorocarbons and hydrofluorocarbons are set to be phased out in  
11 the coming years [3]. From the perspective of an end-user, technical solutions are required that are both  
12 environmentally friendly and economically feasible. This demands the identification of both novel working  
13 fluids that meet all legislated requirements, and ORC systems that are optimised in terms of performance  
14 indicators such as the net-present value or the levelised cost of energy.

15 During a conventional working-fluid selection study an optimal working fluid is typically selected after  
16 screening a group of fluids based on predefined criteria and then conducting a parametric optimisation study  
17 [4, 5]. However, such an approach cannot be used to identify new and potentially novel working fluids, and  
18 therefore more holistic approaches are required. For example, Drescher and Brüggemann [6] identified five  
19 optimal working fluids for a biomass application from an initial group of 1,800 substances, whilst Schwöbel  
20 et al. [7] devised a working-fluid screening process and applied it to 3,174 potential working fluids. More  
21 recently, Preißinger et al. [8] combined computational chemistry techniques with a thermodynamic process  
22 simulation, and applied a multi-criteria evaluation technique to 72 million chemical substances. Other  
23 authors have attempted a more generalised approach to working-fluid selection, by developing correlations  
24 that relate working-fluid parameters, such as the critical temperature, to the heat source conditions [9, 10].

25 Alternatively, computer-aided molecular design (CAMD) can be used to provide a more holistic approach  
26 to working-fluid selection. In CAMD, a potential working fluid is described by functional groups, which can  
27 be put together in different ways to form different molecules. For example, the  $-\text{CH}_3$ ,  $-\text{CH}_2-$ ,  $>\text{C}-$ ,  $>\text{C}<$ ,  
28  $=\text{CH}_2$ , and  $=\text{CH}-$  groups can be combined to generate a large number of hydrocarbon working fluids. Then,  
29 if an equation of state is available that can predict thermodynamic properties based on the functional groups  
30 from which it is composed, the molecular structure of the working fluid can be simultaneously optimised  
31 alongside the ORC system. In this sense, CAMD-ORC models have the potential to identify novel working-  
32 fluids which may otherwise be overlooked, whilst removing preemptive and subjective screening criteria.

33 Papadopoulos et al. [11] used CAMD to identify potential working-fluid candidates before completing  
34 a more conventional ORC process simulation, and later applied CAMD to the optimal design of working-  
35 fluid mixtures [12]. Brignoli and Brown [13] used group-contribution methods to investigate the effect of a  
36 working-fluid's critical point on the thermodynamic performance of the ORC, whilst Palma-Flores et al. [14]  
37 demonstrated the potential of CAMD to identify new fluids with higher thermal efficiencies and better safety  
38 characteristics. Furthermore, Su and Deng [15] developed a thermodynamic ORC model, and later imple-  
39 mented this into a CAMD-ORC framework [16]. Cignitti et al. [17] also developed a CAMD-ORC model,  
40 and in addition to optimising the thermodynamic performance, also considered the heat-exchanger require-

41 ments. However, all of these previous studies have relied on empirical group-contribution methods, such  
42 as the Joback and Reid method [18], to obtain working-fluid parameters including the critical temperature  
43 and pressure from which thermodynamic properties can be calculated using a cubic equation of state. More  
44 advanced group-contribution equations of state have also been applied within a CAMD-ORC framework,  
45 which use molecular-based equations of state based on statistical associating fluid theory (SAFT) [19, 20].  
46 Lampe et al. [21, 22] developed a CAMD-ORC model based on the PC-SAFT equation of state [23, 24],  
47 and used this model to optimise ORC systems for a geothermal application. The CAMD-ORC optimisation  
48 was split into two stages. In the first stage an optimal, but hypothetical, working fluid was identified, and  
49 in the second stage real working fluids with similar performance were identified. Later, Schilling et al. [25]  
50 reduced the problem to a single stage optimisation in which the working-fluid structure and ORC system  
51 are simultaneously optimised, and has recently extended the model to include transport properties and cost  
52 correlations, facilitating the specific-investment cost to be determined [26].

53 With a few exceptions, the major limitation of previous CAMD-ORC models has been a focus on opti-  
54 mising the cycle and its thermodynamic performance; however, achieving the successful commercialisation  
55 of ORC systems across a range of applications requires a consideration of thermoeconomic performance.  
56 Quoilin et al. [27] evaluated the specific-investment cost (SIC) of small-scale waste-heat driven ORC units,  
57 whilst Lecompte et al. [28] optimised the design of ORC units for large-scale CHP plants and waste-heat  
58 recovery. Multi-objective optimisation studies can be also found in the literature [29–31], where the authors  
59 considered the trade-off between maximising power output whilst minimising the SIC. However, all of these  
60 previous thermoeconomic studies consider only predefined working fluids, and conduct a separate optimisa-  
61 tion for each specific fluid. On the contrary, thermoeconomic methods have not been previously applied to  
62 CAMD-ORC models, partly due to the requirement of group-contribution methods for determining transport  
63 properties to size the system heat exchangers.

64 Another limitation to previous CAMD-ORC models is that they typically focus only on a basic, non-  
65 recuperated, subcritical ORC system. However, there exist opportunities to improve the thermodynamic  
66 performance of this basic ORC by changing the cycle architecture. For example, using a working-fluid  
67 mixture instead of a pure fluid results in non-isothermal, isobaric phase change processes, which facilitates a  
68 better thermal match between the working fluid and heat source, and between the working fluid and heat sink,  
69 thus reducing irreversibilities and improving the the thermodynamic performance of the cycle [29, 32, 33].  
70 Alternatively, operating a partially-evaporated cycle, in which expansion occurs from a two-phase state, can  
71 also be used to increase the power output from the system [34, 35].

72 The authors of the current paper have previously developed a CAMD-ORC framework, based on the  
73 SAFT- $\gamma$  Mie group-contribution equation of state [36]. In this previous work, empirical group-contribution  
74 transport property prediction methods for hydrocarbon working fluids were validated against NIST REF-  
75 PROP [37]. The aim of the current study is to combine these transport property prediction methods with  
76 a heat-exchanger sizing model and integrate this model into the CAMD-ORC framework. This, in turn, al-

77 lows a thermoeconomic assessment of the system to be conducted following a thermodynamic optimisation,  
78 and therefore optimal working fluids to be identified based on thermoeconomic performance indicators. In  
79 addition, the current paper extends the thermodynamic model, such that recuperated, partially-evaporated  
80 and working-fluid mixture cycles can all be considered within the CAMD-ORC framework. It is noted that  
81 within this paper the models for the novel cycles are presented and validated, but are not considered in the  
82 case study as suitable methods for predicting heat-transfer coefficients for mixtures, and cost correlations  
83 for two-phase expanders are not available. However, their inclusion is justified as it facilitates these novel  
84 systems to be evaluated in the future. So far as the authors are aware, this is the first study that details a  
85 complete CAMD-ORC framework, based on an advanced group-contribution equation of state, that can  
86 conduct a thermoeconomic assessment in this manner, in addition to simulating novel cycle architectures.

87 In Section 2 the key aspects of the CAMD-ORC framework are discussed, including the group-contribution  
88 methods, the thermodynamic model and the component sizing models, which are then validated in Section  
89 3. In Section 4 the framework is applied to a case study considering the design of hydrocarbon working  
90 fluids. Finally, the key findings from this study are discussed in Section 6.

## 91 **2. CAMD-ORC model**

### 92 *2.1. Group-contribution methods*

93 Group-contribution methods determine the properties of a particular molecule based on the functional groups  
94 that make it up. For example, isopentane is described by three  $-\text{CH}_3$  groups, one  $-\text{CH}_2-$  and one  $>\text{CH}-$   
95 group. In a group-contribution method group parameters are only required for the individual groups, which  
96 allows the evaluation of novel working fluids for which property prediction would not be possible using  
97 conventional approaches. To capture the trade-off between thermodynamic performance and system costs,  
98 group-contribution methods are required for both the thermodynamic properties and transport properties.  
99 In this work, the SAFT- $\gamma$  Mie equation of state [38] is used for thermodynamic property prediction. SAFT- $\gamma$   
100 Mie is a state-of-the-art version of statistical associating fluid theory (SAFT) [19, 20] wherein a Mie potential  
101 is used to model the interaction between two molecular groups [38]. Group parameters are available for  
102 the hydrocarbon groups considered within this paper, and have been validated against experimental data  
103 [39]. Unfortunately, SAFT- $\gamma$  Mie is only suitable for determining thermodynamic properties, so alternative  
104 methods are required to predict the dynamic viscosity  $\mu$ , thermal conductivity  $k$  and surface tension  $\sigma$ .  
105 Previously, empirical group-contribution methods for the prediction of these properties have been applied to  
106 hydrocarbon working fluids, and validated against data from NIST [36]. The correlations applied here are  
107 summarised in Table 1, and are reviewed in detail in Ref. [36].

### 108 *2.2. Thermodynamic modelling*

109 The thermodynamic analysis of the ORC is well described within the literature, and consists of applying an  
110 energy balance to each component within the cycle. Besides analysing a sub-critical, non-recuperated cycle,

Table 1: Summary of group-contribution methods used within the CAMD-ORC framework.

Property	Liquid phase	Vapour phase
Thermodynamic ( $T, p, h, s, \rho$ )	SAFT- $\gamma$ Mie [38]	
Critical ( $T_{cr}, p_{cr}, V_{cr}$ )	Joback-Reid [18]	
Surface tension ( $\sigma$ )	Sastri-Rao [40]	
Dynamic viscosity ( $\mu$ )	Joback-Reid [18] ( $n$ -alkanes)	Reichenberg [41, 42]
	Sastri-Rao [43] (branched alkanes)	
Thermal conductivity ( $k$ )	Sastri [44]	Chung [45, 46]

111 the CAMD-ORC model has also been extended to be suitable for the evaluation of cycles operating with  
 112 mixtures, recuperated cycles, and cycles with partial evaporation. A schematic representation of each cycle  
 113 architecture, and the prescribed notation is given in Figure 1.

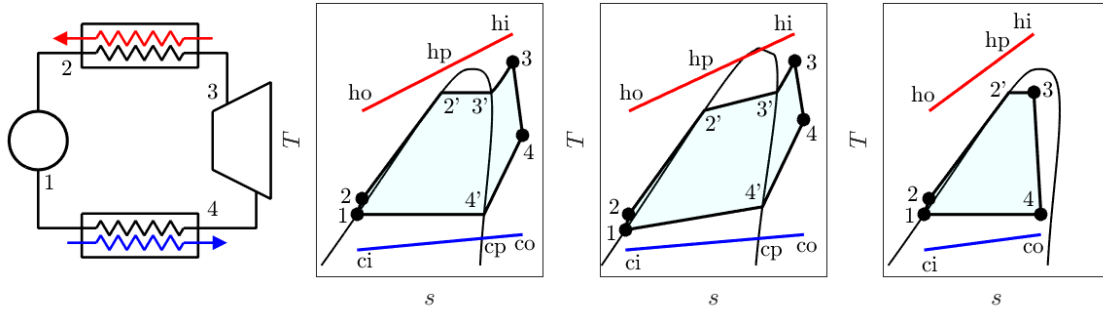


Figure 1: Schematic of the ORC system and the different cycle architectures represented on a  $T$ - $s$  diagram. From left to right: cycle schematic, basic non-recuperated, mixture and partially-evaporated.

114 For all cycles, the system is assumed to be in a steady state, pressure drops within the heat exchangers  
 115 and piping are neglected, whilst the condensation temperature  $T_1$  and reduced pressure  $p_r$  ( $p_2/p_{cr}$ , where  $p_2$   
 116 and  $p_{cr}$  are the evaporation and critical pressures respectively) are both defined as optimisation variables.  
 117 Moreover, values for the pump isentropic efficiency  $\eta_p$  and expander isentropic efficiency  $\eta_e$  are fixed. Within  
 118 this paper, the expander is assumed to be radial turbine, which is capable of achieving a large expansion  
 119 ratio across a single stage, and is suitable for the power range being considered. The authors rightfully  
 120 acknowledge that assuming a single fixed turbine isentropic efficiency for a range of system sizes operating  
 121 with different expansion ratios is an oversimplification. However, these effects have been neglected owing to  
 122 the complexity of requiring a more detailed expander model, which is not a particular focus of this work,  
 123 but could be easily included in the future.

Alongside  $T_1$  and  $p_r$  an additional optimisation variable is required to define all the state points within  
 the cycle, and this is defined using the notation  $z$ . This parameter is introduced to allow both superheated  
 and partially-evaporated cycles to be modelled using one optimisation variable, which varies between 0 and

2. When  $0 \leq z \leq 1$ , two-phase expansion is assumed and  $z$  is equal to the expander inlet vapour quality. When  $1 < z \leq 2$ , the working fluid expands from a superheated state and the amount of superheating  $\Delta T_{\text{sh}}$  is given by:

$$\Delta T_{\text{sh}} = (z - 1)(T_{\text{hi}} - T_{3'}), \quad (1)$$

124 where  $T_{\text{hi}}$  is the heat-source inlet temperature and  $T_{3'}$  is the saturated-vapour temperature. It is noted  
 125 that when  $z = 2$ ,  $T_{\text{hi}} = T_3$  which would imply an infinitely large heat exchanger. Therefore, within an  
 126 optimisation a minimum evaporator pinch constraint is imposed ( $T_{\text{hi}} - T_{3'} > PP_{\text{h,min}}$ ), which will always  
 127 result in cycles where  $z < 2$ .

The working-fluid mass flow rate  $\dot{m}_o$  (kg/s) is determined by imposing the evaporator pinch point  $PP_{\text{h}}$  at the start of evaporation (*i.e.*,  $PP_{\text{h}} = T_{\text{hp}} - T_{2'}$ ), and applying an energy balance:

$$\dot{m}_o = \frac{(\dot{m}c_p)_{\text{h}}(T_{\text{hi}} - T_{\text{hp}})}{h_3 - h_{2'}}, \quad (2)$$

where  $(\dot{m}c_p)_{\text{h}}$  is the heat-source heat-capacity rate (W/K) and  $h_{2'}$  and  $h_3$  are the enthalpies (J/kg) of the working fluid at the start of evaporation and expander inlet respectively. With the mass-flow rate known, the thermodynamic performance of the ORC can be evaluated by determining the net power output from the system  $\dot{W}_{\text{n}}$  (W):

$$\dot{W}_{\text{n}} = \dot{m}_o [(h_3 - h_4) - (h_2 - h_1)]. \quad (3)$$

Finally, an energy balance is applied to the condenser to obtain the condenser pinch point  $PP_{\text{c}}$ . This is given by:

$$PP_{\text{c}} = T_4 - \frac{\dot{m}_o(h_4 - h_1)}{(\dot{m}c_p)_{\text{c}}}, \quad (4)$$

if the expansion process ends in two-phase region, and:

$$PP_{\text{c}} = T_{4'} - \frac{\dot{m}_o(h_{4'} - h_1)}{(\dot{m}c_p)_{\text{c}}}, \quad (5)$$

128 if the expansion process ends in the superheated region. Within the model, the minimum allowable condenser  
 129 pinch point  $PP_{\text{c,min}}$  is defined as a constraint.

130 The calculation process described so far is applicable to all types of cycle, but for working-fluid mix-  
 131 tures and recuperated cycles, additional parameters are introduced. For a mixture both working fluids are  
 132 described by their functional groups and the variable  $x$  is introduced to represent the mass fraction of the  
 133 first fluid. A recuperated cycle is modelled by a fixed recuperator effectiveness  $\epsilon_{\text{r}}$ , and the inclusion of a  
 134 recuperator is defined by a binary flag.

### 135 2.3. Component sizing

136 The evaporator and condenser are assumed to be tube-in-tube heat exchangers, which are cost effective for  
 137 small- to medium-scale applications [47]. The heat exchangers are sized by determining the total required  
 138 heat-transfer area, which is obtained by calculating the heat-transfer coefficient in the different single- and  
 139 two-phase heat-transfer regions. In the evaporator this corresponds to single-phase preheating, two-phase



140 evaporation and single-phase superheating regions, and in the condenser it corresponds to single-phase  
 141 desuperheating and two-phase condensation regions.

142 Depending on the heat-transfer region, different Nusselt-number correlations are applied to determine the  
 143 local heat-transfer coefficient. For single-phase heat transfer the Dittus-Boelter [48] correlation has been used.  
 144 For evaporation, the correlations proposed by Cooper [47] and Gorenflo [49] have been used for nucleate-  
 145 boiling conditions, whereas the Dobson [50] and Zuber [47] correlations have been used to account for the  
 146 convective-heat-transfer phenomena. For condensation inside tubes, the correlations proposed by Shah [51]  
 147 and Dobson [50] have been considered, accounting for both gravity-driven and shear-driven condensation.  
 148 The reader can refer to Ref. [52] for a detailed analysis and comparison of the correlations selected. The  
 149 Nusselt-number correlations for two-phase heat transfer are typically a function of the vapour quality, which  
 150 varies along the length of the heat exchanger. Therefore, the heat-exchanger length is discretised into  $n$   
 151 segments of equal heat duty. For each segment the vapour quality is assumed to be constant and an estimate  
 152 for the heat-transfer area for that segment is obtained.

Expressed mathematically, the total heat-transfer area for a given heat exchanger  $A$  ( $\text{m}^2$ ) is given as the summation of all the segments:

$$A = \sum_{i=1}^n \frac{\dot{Q}_i}{U_i \Delta T_{\log,i}}, \quad (6)$$

153 where  $\dot{Q}_i$  and  $\Delta T_{\log,i}$  are the heat-transfer rate (W) and counter-flow log-mean temperature difference (K)  
 154 for segment  $i$  respectively, and  $U_i$  is the overall heat-transfer coefficient ( $\text{W}/(\text{m}^2 \text{K})$ ) for segment  $i$  and is  
 155 found based on the heat-transfer coefficients either side of heat-exchanger wall.

A key consideration when estimating the required heat-transfer area is the pressure drop along the full length of the heat exchanger. In this study, a number of pressure drop correlations have been used to predict the pressure drop of the organic working fluid, the heat source fluid (Therminol 66) and the cooling fluid (water) inside the heat exchangers. For both the evaporator and the condenser units the pressure drop was restricted to not exceed 1–2 bar, which is in line with good-practice industry standards. For the single-phase zone, the pressure drop is calculated as a function of the fluid velocity inside the tubes, the diameter of tubes, the length of the heat exchanger, and a friction coefficient. The calculation is completed using the following set of equations in line with [47]:

$$Re = \rho u D / \mu; \quad (7)$$

$$f = 0.046 Re^{-0.2}; \quad (8)$$

$$\Delta P = 4f \frac{L}{D} \frac{\rho u^2}{2}, \quad (9)$$

156 where  $Re$  is the Reynolds number,  $\rho$  is the fluid density ( $\text{kg}/\text{m}^3$ ),  $u$  is the fluid velocity (m/s),  $D$  is the tube  
 157 diameter (characteristic length) (m),  $\mu$  is the dynamic viscosity (Pa s),  $f$  is a friction factor (also referred to  
 158 as the Fanning friction factor),  $L$  is the heat exchanger length (m), and  $\Delta P$  is the pressure drop (Pa). For

159 the two-phase zone pressure drop, the correlation developed by Chisholm as presented in Ref. [47] has been  
 160 used. The equations for the two phase zone pressure drop have been omitted here for brevity.

161 *2.4. Thermoeconomic analysis*

Since there are only a limited number of ORC applications worldwide, and system cost data are not publicly available, cost correlations originating from the chemical industry are commonly used in the literature. A well-established method is the module costing technique [53], which provides the costs of individual components, based on a specific sizing attribute (*e.g.*, heat-transfer area for heat exchangers *etc.*). By adding the individual component costs the total ORC unit cost is obtained. The costing method applied within this study is summarised in Ref. [52] and uses the cost correlations given by Seider et al. [54]:

$$C_p^0 = F \exp(Z_1 + Z_2 \ln(X) + Z_3 \ln(X)^2 + Z_4 \ln(X)^3 + Z_5 \ln(X)^4), \quad (10)$$

and Turton et al. [55]:

$$C_p^0 = F10^{(Z_1+Z_2 \log(X)+Z_3 \log(X)^2)}, \quad (11)$$

162 where  $C_p^0$  is the component cost in £;  $F$  is a material factor accounting for the component manufacturing;  $Z_i$   
 163 is the cost coefficient; and  $X$  is the sizing attribute. Both  $Z_i$  and  $X$  vary depend on the type of the equipment  
 164 selected and the values used to estimate the purchase cost of each piece of equipment are summarised in  
 165 Table 2. It is assumed that the pump is a centrifugal pump, whilst the heat exchangers are of tube-in-tube  
 166 construction. As previously stated, the expander is assumed to be a radial turbine, and this component's  
 167 cost is based only on the power output. In reality, the expansion ratio of the turbine will impact both the  
 168 expander efficiency and cost. Within this work these effects have been neglected owing to the complexities  
 169 of requiring a more detailed expander model, and because correlations that consider these effects are either  
 170 not currently available, or not sufficiently validated. However, these effects should be considered in future  
 171 studies. Finally, the Chemical Engineering Plant Cost Index (CEPCI) is used to convert the cost to today's  
 172 values. For Turton et al. [55] the basis year is 2001 ( $CEPCI_{2001} = 397$ ), whilst for Seider et al. [54] the basis  
 173 year is 2006 ( $CEPCI_{2006} = 500$ ). The costs are converted to today's values using  $CEPCI_{2017} = 562.1$ .

Table 2: Cost correlations coefficients

Component	Attribute ( $X$ )	$F$	$Z_1$	$Z_2$	$Z_3$	$Z_4$	$Z_5$	Ref.
Expander	Power, $\dot{W}_e$ (kW)	3.5	2.2486	1.4965	-0.1618	0	0	[55]
Pump	$S^*$	2.7	9.2951	-0.6019	0.0519	0	0	[54]
Pump motor	Power, $\dot{W}_p$ (HP)	1.4	5.83	0.134	0.0533	0.0286	0.00355	[54]
Evaporator - Condenser	Area ( $m^2$ )	1	9.5638	0.532	-0.0002	0	0	[54]
Preheater - Desuperheater	Area ( $m^2$ )	1	10.106	-0.4429	0.0901	0	0	[54]

\*  $S = \dot{V}\sqrt{H}$  where  $\dot{V}$  is the pump volumetric flow rate in gallons per minute and  $H$  is the pump head in feet.

The CAMD-ORC framework is formulated in gPROMS [56], and the optimisation is completed using the OAERAP outer-approximation algorithm. The optimisation concerns integer variables describing the working-fluid molecular structure and continuous variables describing the power system, and therefore is a mixed-integer non-linear programming (MINLP) problem. The optimisation is solved by first relaxing the integer variables to continuous variables and completing a non-linear programming (NLP) optimisation, which in turn supplies a maximum for the objective function. The MINLP is then solved by successive iterations of a mixed-integer linear programming problem (MILP), in which the objective function and constraints are linearised, and an additional NLP in which the power system variables are optimised for a particular fluid identified from the MILP. The general optimisation is therefore formulated as:

$$\max f(\mathbf{x}, \mathbf{y}), \quad (12)$$

subject to:

$$\mathbf{x}_{\min} \leq \mathbf{x} \leq \mathbf{x}_{\max}; \quad (13)$$

$$\mathbf{y}_{\min} \leq \mathbf{y} \leq \mathbf{y}_{\max}; \quad (14)$$

$$g(\mathbf{x}, \mathbf{y}) \leq 0; \quad (15)$$

$$h(\mathbf{x}, \mathbf{y}) \leq 0; \quad (16)$$

where  $f(\mathbf{x}, \mathbf{y})$  is the objective function to be maximised,  $\mathbf{x}$  and  $\mathbf{y}$  are vectors containing system and working-fluid variables respectively, inequalities 13 and 14 represent the lower and upper bounds for the variables, and  $g(\mathbf{x}, \mathbf{y})$  and  $h(\mathbf{x}, \mathbf{y})$  are the cycle and molecular constraints respectively.

Within this study, the objective of the optimisation is to maximise power output  $\dot{W}_n$  and then assess these optimal systems from an economic perspective. However, future research should integrate the economic analysis into the optimisation model, and facilitate technoeconomic performance indicators, such as specific-investment cost, to be considered. Such an optimisation could be easily carried out in the future using the existing CAMD-ORC framework.

### 3. Model validation

#### 3.1. Thermodynamic and transport property validation

Within the CAMD-ORC framework group-contribution methods are used to predict both the thermodynamic and transport properties of the working fluid. As discussed previously, SAFT- $\gamma$  Mie is used to predict the thermodynamic properties, and an array of different empirical correlations are used for the transport properties. The non-group-contribution formulation of SAFT- $\gamma$  Mie, SAFT-VR Mie [57], has previously been applied to the study of optimal working-fluid mixtures for ORC systems, in which the average absolute deviation in saturation properties (density and pressure), specific-heat capacities and critical properties

191 (temperature and pressure), in comparison to the values provided by NIST REFPROP, are all below 5% for  
192 the pure alkane and perfluoroalkane fluids considered [58]. Furthermore, SAFT- $\gamma$  Mie has also been shown  
193 to provide an accurate description of fluid-phase thermodynamic properties for a wider variety of fluids  
194 [37–39, 59]. Moreover, the authors of the current paper have also validated SAFT- $\gamma$  Mie within the context  
195 of the existing CAMD-ORC framework [36], which also involved the validation of the group-contribution  
196 transport property prediction methods. It was found that the absolute deviations of the critical temperatures  
197 and pressures of the selected hydrocarbon working fluids were less than 0.5% and 4% respectively. The  
198 average absolute deviations of the viscosity, thermal conductivity and surface tension with respect to available  
199 experimental data, evaluated at temperatures between 20 °C and 400 °C, were generally less than 4%, 5%  
200 and 8% respectively. Overall, this provides reasonable confidence in the suitability of the group-contribution  
201 methods employed within this current work.

### 202 3.2. Cycle modelling

203 In our previous work, the CAMD-ORC model has been validated for a non-recuperated, sub-critical ORC  
204 operating with an array of hydrocarbon working fluids, by comparing the results to a similar model that  
205 uses the NIST REFPROP program to calculate thermodynamic properties [36]. Both models were found to  
206 identify optimal cycles with very similar reduced evaporating pressures, whilst the maximum power output  
207 predicted by the CAMD-ORC model deviated by less than 1.5% compared to the REFPROP model.

208 In this section, a similar validation study will be completed to confirm the suitability of the CAMD-ORC  
209 model to simulate the alternative cycle architectures that were introduced in Section 2.2, namely cycles  
210 operating with working-fluid mixtures and partially-evaporated cycles. Although the model is expected to  
211 be applicable for the range of heat-source temperatures relevant to ORC systems (*i.e.*, 373–673 K), for the  
212 validation study the heat source is assumed to be at 473 K. It is easily shown that the thermodynamic  
213 performance of an ORC system is independent of the heat-source mass-flow rate and therefore the heat  
214 source is defined with a heat-capacity rate of  $\dot{m}c_p = 4.2$  kW/K. Finally, the pump and expander are  
215 modelled assuming fixed isentropic efficiencies of  $\eta_p = 0.7$  and  $\eta_e = 0.8$  respectively, whilst  $T_1 = 303$  K and  
216  $PP_{h,\min} = 10$  K are assumed; these values considered to be representative of a typical ORC system.

#### 217 3.2.1. Partially-evaporated cycles

218 The purpose of the first validation study is to confirm the suitability of the CAMD-ORC model for partially-  
219 evaporated cycles. For this study, five different working fluids have been considered, namely *n*-pentane,  
220 *n*-hexane, *n*-heptane, isopentane and isohexane. For these five fluids a parametric study was completed in  
221 which  $z$  was varied between 0 and 2, and this was repeated at different evaporation pressures. A comparison  
222 between the results obtained using the CAMD-ORC model, based on SAFT- $\gamma$  Mie, and a similar model using  
223 NIST REFPROP is given in Figure 2. In this figure, the results for only three fluids are shown, however  
224 the other two fluids were found to follow the same behaviour. The 18.6 and 24.9 bar cases for *n*-hexane do

not appear in Figure 2 because the saturation temperature of  $n$ -hexane at these pressures is higher than the heat-source temperature.

Overall, it is observed that, except for the 28.0 bar cases for  $n$ -pentane and isopentane, a very good agreement between the two models is obtained. Neglecting these two cases, the maximum deviations between the CAMD-ORC model and REFPROP model are below 4%, 1% and 5% for  $n$ -pentane,  $n$ -hexane and isopentane respectively. For  $n$ -heptane and isohexane, the maximum deviations are below 2% and 3% respectively. Ultimately, this validates the suitability of the CAMD-ORC model to simulate these types of cycles. The deviation for the 28.0 bar isopentane case can be explained because at higher pressures the cycle is operating closer to the critical point (33.8 bar), and the deviation between SAFT- $\gamma$  Mie and NIST REFPROP is found to increase as the critical point is approached.

More generally, from Figure 2 it is observed that the maximum power is always generated when  $z < 1$ . This is due to the fact that expansion when  $z < 1$  takes place inside the two-phase region, such that a larger proportion of the heat transfer during heat addition occurs during preheating, and results in a lower heat-source outlet temperature,  $T_{ho}$ . This, in turn, means that the ORC is capable of extracting more heat from the heat source, leading to a higher power output. This clearly indicates the potential thermodynamic performance benefit of allowing the working fluid to expand from a two-phase state.

### 3.2.2. Working-fluid mixtures

For the validation of the CAMD-ORC model for mixtures, three different mixtures have been considered, namely  $n$ -hexane/ $n$ -butane,  $n$ -heptane/ $n$ -butane and  $n$ -heptane/ $n$ -pentane. For each mixture a parametric study was completed in which the mass fraction of Fluid 1,  $x$ , was varied from 0 to 1, and this was repeated at different evaporation pressures. In each case, it was assumed that expansion occurs from a saturated-vapour state (i.e.,  $z = 1$ ). The comparison between the results obtained using the CAMD-ORC model, and the model based on NIST REFPROP is shown in Figure 3. Again, a very good agreement is observed for the different case studies; neglecting power outputs below 20 kW, which do not represent good thermodynamic cycles, the maximum deviations between the CAMD-ORC and REFPROP models are 4%, 4% and 8% for the  $n$ -hexane/ $n$ -butane,  $n$ -heptane/ $n$ -butane and  $n$ -heptane/ $n$ -pentane cases respectively. These results therefore validate the CAMD-ORC model for simulating these types of cycles.

### 3.3. Heat-exchanger sizing validation

The CAMD-ORC framework has previously been used to optimise the working fluid and cycle conditions for a non-recuperated, basic ORC, and this process was completed for three different waste-heat streams [36]. Now, using the group-contribution transport-property prediction methods and the heat-exchanger sizing model described in Sections 2.1 and 2.3 respectively, it is possible to determine the area requirements of the evaporator and the condenser for these optimum cycles. The full analysis will be described in detail in Section 4, however, first, it is necessary to validate the developed model.

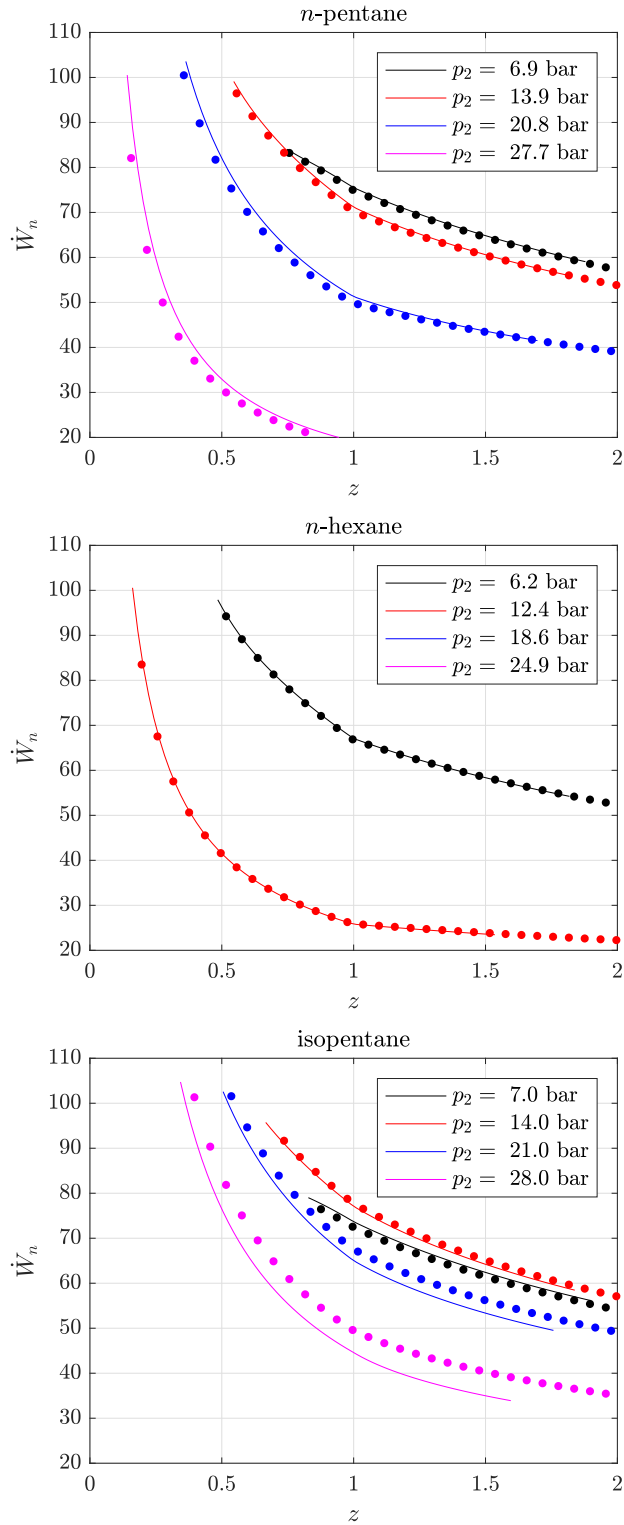


Figure 2: Comparison between the power output ( $\dot{W}_n$  in kW) predicted by the CAMD-ORC model (circular markers) and predicted by a model using NIST REFPROP for thermodynamic properties (continuous curves) for three different working fluids operating within a partially-evaporated ( $z < 1$ ) and a superheated ( $z \geq 1$ ) cycle.

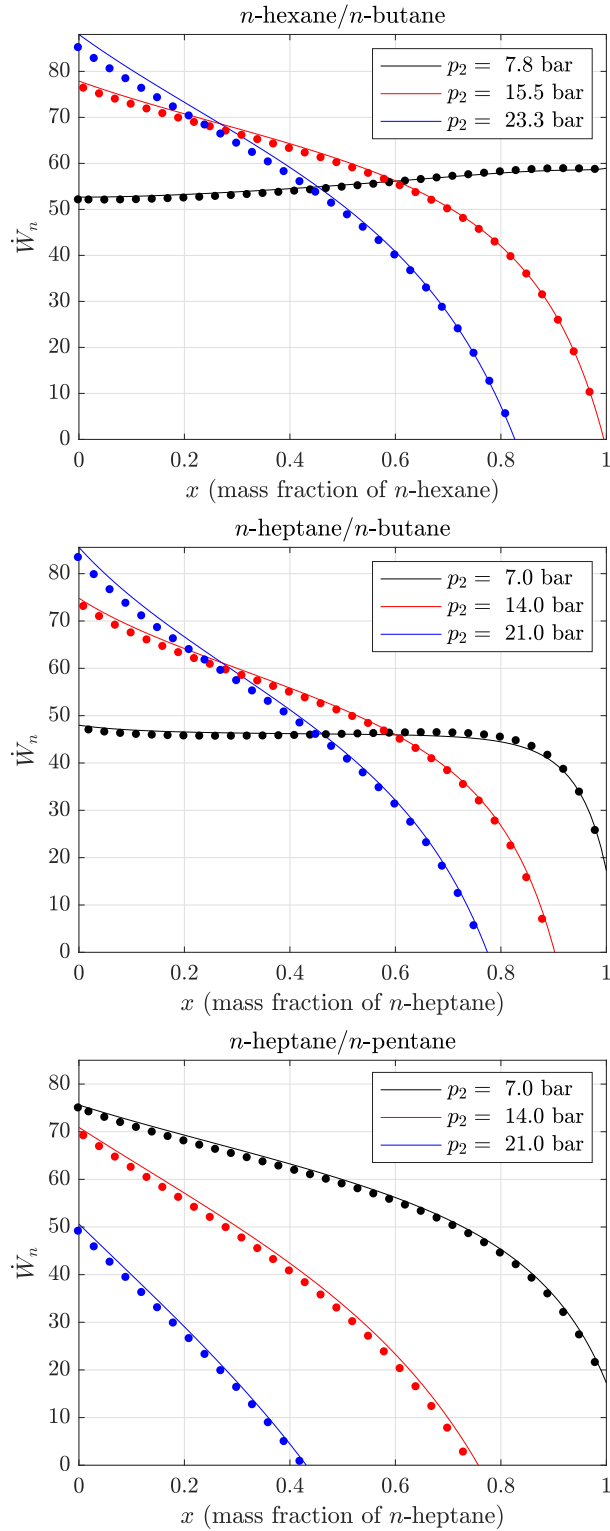


Figure 3: Comparison between the power output ( $\dot{W}_n$  in kW) predicted by the CAMD-ORC model (circular markers) and predicted by a model using NIST REFPROP for thermodynamic properties (continuous curves) for three different fluid mixtures.

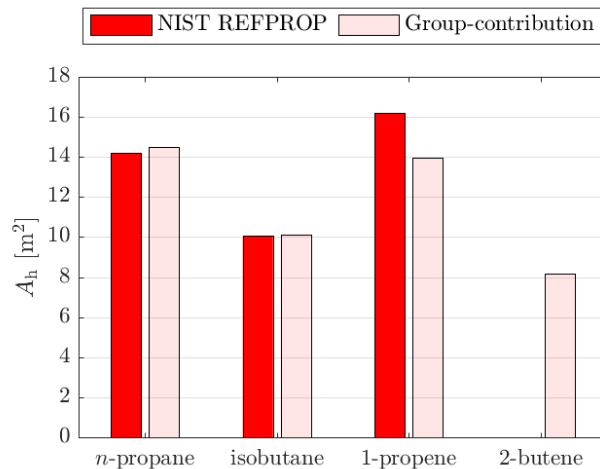


Figure 4: Comparison between the evaporator area ( $A_h$  in  $m^2$ ) obtained using group-contribution transport properties and NIST REFPROP.

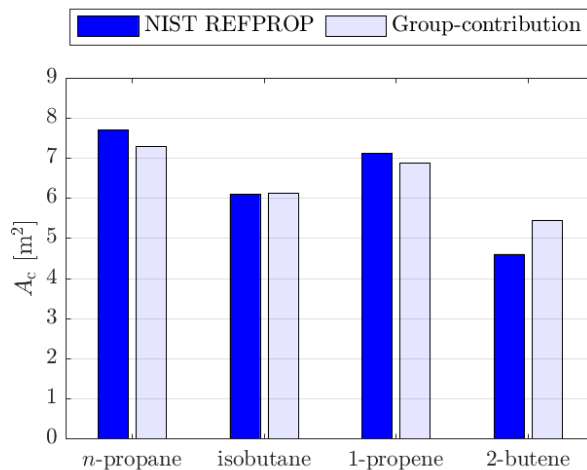


Figure 5: Comparison between the condenser area ( $A_c$  in  $m^2$ ) obtained using group-contribution transport properties and NIST REFPROP.

259 For this validation study, the heat-exchanger sizing is first performed using properties obtained from  
 260 NIST REFPROP. Then, these results are compared to those obtained when the group contribution transport  
 261 properties are used. Not all of the fluids considered within the initial CAMD-ORC optimisation study are  
 262 available within REFPROP, and therefore it is only possible to validate the model for a subset of the fluids  
 263 considered. This subset of fluids used for the comparison study includes *n*-propane (*n*-alkane), isobutane  
 264 (methyl alkanes), 1-propene (1-alkene) and cis-2-butene (2-alkene). The heat-carrier fluid for all fluids is  
 265 Therminol 66<sup>®</sup>, entering the evaporator at 150 °C and 1 bar.

266 In Figures 4 and 5, the evaporator and condenser area requirements for the four working fluids are pre-



267 sented. In line with the figures, the results obtained using the group-contribution transport property model  
268 are in good agreement with those obtained from NIST REFPROP. The heat-exchanger area calculations  
269 for *n*-propane and isobutane have negligible difference between the two methods. The highest deviation is  
270 recorded for *cis*-2-butene, where the condenser unit surface area is overestimated by the group-contribution  
271 method by approximately 18%, being on the conservative side of the heat-exchanger design. It should be  
272 noted that the Nusselt number correlations for the evaporator area calculation require the use of the working  
273 fluid surface tension, which for *cis*-2-butene is not available in NIST.

274 Compared to the *n*-propane and isobutane cases, the 1-propene case shows a relatively large deviation of  
275 13.9% between the total evaporator area predicted using the CAMD-ORC model and using REFPROP. This  
276 deviation occurs, in part, because it is not possible to match exactly the thermodynamic cycle conditions  
277 input into the heat-exchanger sizing model, and those output from the CAMD-ORC model. This, coupled  
278 with the 1-propene cycle having a higher degree of superheat, which is 54 °C compared to 18 °C and  
279 0.1 °C for *n*-propane and isobutane cycles respectively, results in the 1-propene superheater load for the  
280 REFPROP heat-exchanger sizing model being higher than the CAMD-ORC model. Considering that the  
281 heat-transfer coefficient for a vapour is significantly lower than for a liquid or two-phase fluid, this results in  
282 the REFPROP model predicting a superheater area of 4.37 m<sup>2</sup>, compared to 3.34 m<sup>2</sup> for the CAMD-ORC  
283 model, thus accounting for the relatively large deviation observed.

284 Ultimately, the deviation introduced by inputting the cycle parameters from the CAMD-ORC model into  
285 the REFPROP heat-exchanger sizing model does not directly represent an issue with the group-contribution  
286 transport property prediction methods but is instead a carry-over from the difference between the ther-  
287 modynamic properties predicted by SAFT- $\gamma$  Mie within the thermodynamic cycle model and REFPROP.  
288 Having said this, neglecting the 2-butene condenser and 1-propene evaporator, the percentage deviation  
289 between the CAMD-ORC model and REFPROP for the remaining heat exchangers are all below 5%. These  
290 values are very much in line with the percentage deviations observed between the thermodynamic properties  
291 (< 5%), and transport properties (< 8%) discussed in Section 3.1. Overall, this gives good confidence in the  
292 heat-exchanger sizing model implemented within the CAMD-ORC model.

#### 293 4. Case study

294 As already discussed, the CAMD-ORC framework has previously been used to optimise the working fluid and  
295 thermodynamic cycle for three different waste-heat streams [36]. Furthermore, the transport-property group-  
296 contribution correlations have been coupled to the heat-exchanger sizing model, and the heat-transfer area  
297 requirements for a few of the optimal cycles that resulted from the initial thermodynamic study have been  
298 determined in the previous section. The aim of this case study is to determine the heat-transfer requirements  
299 for a larger group of working fluids, and determine the total specific-investment cost (SIC) for each working  
300 fluid. This, in turn, allows optimal cycle configurations to be identified based on thermoeconomics.

301 *4.1. Case study assumptions*

302 It should be noted that within this case study only a basic, non-recuperated ORC has been considered,  
 303 despite the CAMD-ORC being successfully extended to, and validated for, alternative cycle architectures  
 304 within this paper. This is because the uncertainties that are introduced when considered these novel cycle  
 305 architectures. Firstly, sizing the heat exchangers for a cycle operating with a working-fluid mixture would  
 306 require suitable mixing rules to be defined to determine the necessary transport properties. Moreover, there  
 307 also exist large uncertainties in predicting the local heat-transfer coefficient for a working-fluid mixture.  
 308 Secondly, whilst partially evaporated cycles are an extremely interesting idea from the point of view of  
 309 maximising power output, commercial expander technologies for two-phase expansion, with a few exceptions  
 310 [60, 61], are not widely available. Therefore, it follows that cost correlations for two-phase expanders do not  
 311 exist. With this in mind, it follows that a basic, non-recuperated ORC system is the easiest to evaluate from  
 312 a thermoeconomic point of view, and is therefore the best cycle with which to demonstrate the CAMD-ORC  
 313 framework that has been developed within this paper. Nonetheless, as more research into working-fluid  
 314 mixtures and partially-evaporated cycles is conducted, the same tool can be used to evaluate these novel  
 315 cycles, with minimal changes required to the CAMD-ORC framework.

316 Moving back to the case study, the three heat-sources considered are each defined by a heat-capacity rate  
 317 ( $\dot{m}c_p$ ) of 4.2 kW/K, and are defined at 150, 250 and 350 °C respectively. The assumptions for the study are  
 318 listed in Table 3, whilst the working fluids under consideration are given in Table 4. For all three heat-source  
 319 temperatures, the heat source is assumed to be the heat-transfer oil Therminol 66<sup>®</sup> at 1 bar, and the heat  
 320 sink is water. As stated previously, steady-state operating conditions are assumed, and pressure drops within  
 321 the heat exchangers and piping are neglected.

Table 3: Values of the quantities used in the ORC thermodynamic study completed in Ref. [36].

$T_{hi}$ °C	$(\dot{m}c_p)_h$ W/K	$T_{ci}$ °C	$c_{p,c}$ J/(kg K)	$\dot{m}_c$ kg/s	$\eta_p$ %	$\eta_e$ %	$PP_{h,min}$ °C	$PP_{c,min}$ °C	$P_{1,min}$ bar
150, 250, 350	4200	15	4200	5.0	70	80	10	5	0.25

Table 4: Working-fluid groups considered within this study.

<i>n</i> -alkanes	methyl alkanes	1-alkenes	2-alkenes
$\text{CH}_3-(\text{CH}_2)_n-\text{CH}_3$	$(\text{CH}_3)_2-\text{CH}-(\text{CH}_2)_n-\text{CH}_3$	$\text{CH}_2=\text{CH}-(\text{CH}_2)_n-\text{CH}_3$	$\text{CH}_3-\text{CH}=\text{CH}-(\text{CH}_2)_n-\text{CH}_3$

322 The objective of the optimisation is to identify the working fluid and cycle parameters that result in the  
 323 best thermodynamic performance, and then assess these optimal systems from an economic perspective. The  
 324 objective is therefore to maximise power output  $\dot{W}_n$ . For this study, there are five optimisation variables,  
 325 and these are listed in Table 5 alongside the bounds and constraints for the optimisation.

Table 5: Bounds for the optimisation variables and constraints applied during the optimisation.

Variable	Upper bound	Lower bound	Unit	Constraints
$T_1$	288	353	K	$T_{ho} - T_2 \geq PP_{h,min}$
$p_r$	0.001	0.85	-	$T_{hp} - T_{2'} \geq PP_{h,min}$
$z$	1.0	2.0	-	$T_{hi} - T_3 \geq PP_{h,min}$
$PP_h$	10	200	K	$T_1 - T_{cp} \geq PP_{c,min}$
$-(CH_2)_n-$	0	10	-	$T_{4'} - T_{cp} \geq PP_{c,min}$

#### 4.2. Thermodynamic performance

For each working-fluid group in Table 4 a parametric study was completed whereby the number of  $-CH_2-$  groups was varied, and the ORC thermodynamic variables were optimised to maximise the power output from the system [36]. The results from this parametric study are plotted in terms of the power output in Figure 6; here  $C_n$  refers to the number of carbon atoms in the molecule ( $n$ -alkane, methyl alkane, 1-alkene or 2-alkene). The optimal working fluids for the three heat source temperatures of 150, 250 and 350 °C are  $n$ -propane ( $n$ -alkane,  $C_n = 3$ ), 2-pentene (2-alkene,  $C_n = 5$ ) and 2-hexene (2-alkene,  $C_n = 6$ ), corresponding to maximum power outputs of 35.2, 136.7 and 219.0 kW respectively. The corresponding thermal cycle efficiencies are 9.7%, 16.9% and 17.8% respectively.

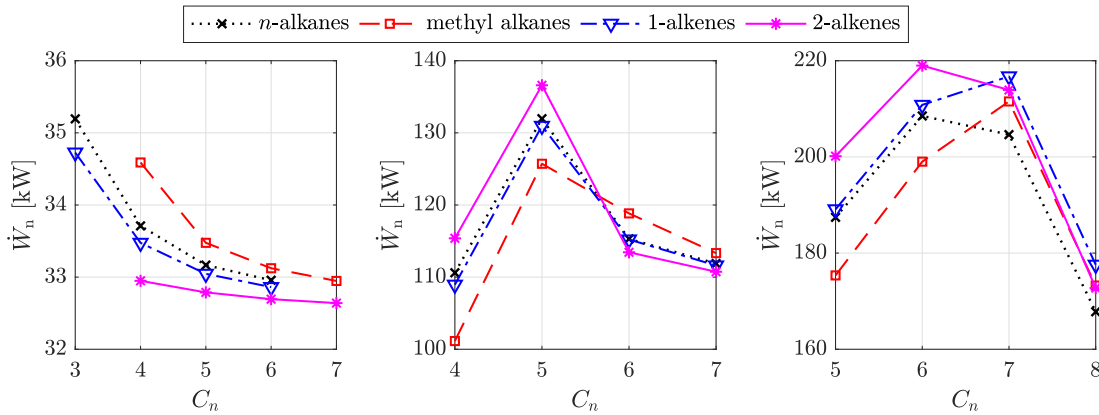


Figure 6: Optimal net power output from an ORC system operating with different hydrocarbon working fluids. Results are plotted against the number of carbon atoms  $C_n$  in the molecule ( $n$ -alkane, methyl alkane, 1-alkene or 2-alkene, as indicated). From left to right:  $T_{hi} = 150, 250, 350$  °C.

The optimal cycles that correspond to the maximum power are explored in Figure 7, in which are displayed three of the cycles ( $C_n = 4, 5$  and 6) for the  $n$ -alkane, 250 °C case-study on a  $T$ - $s$  diagram.

When  $C_n = 4$ , the evaporation temperature, and therefore evaporation pressure, is constrained by the critical temperature since we are only considering subcritical cycles. This results in an optimal cycle with a high reduced pressure and a large amount of superheating, since the high-temperature heat can only be absorbed by the cycle by increasing the working-fluid temperature whilst maintaining the same pressure.

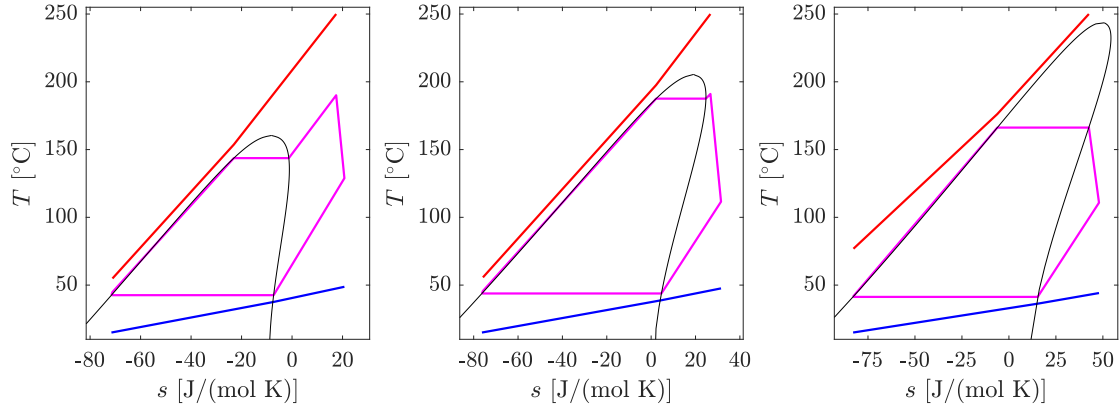


Figure 7:  $T$ - $s$  plots for three cycles from the  $n$ -alkane, 250 °C case-study. From left to right:  $C_n = 4$  ( $n$ -butane), 5 ( $n$ -pentane) and 6 ( $n$ -hexane). The red and blue lines are the heat-source and heat-sink streams, the magenta lines are the ORC and the black is the working-fluid saturation dome.

341 This introduces the need for an additional heat exchanger, namely the superheater, in addition to increasing  
 342 the irreversibilities within the heat-addition process, owing to the increased temperature difference between  
 343 the heat source and working fluid in the evaporation and superheating regions, resulting in a 16% reduction  
 344 in the power output compared to the optimal cycle. It is also noted that the minimum allowable evaporator  
 345 pinch point is observed at the preheater inlet in addition to the evaporator inlet. This corresponds to the the  
 346 lowest heat-source outlet temperature, indicating that the ORC absorbs the maximum amount of heat from  
 347 the available heat source. Arguably, the thermodynamic performance of the  $C_n = 4$  cycle could be improved  
 348 by increasing the evaporation pressure above the critical pressure, and thus operate a transcritical cycle.  
 349 However, it is worth noting that higher evaporation pressures lead to more expensive system components,  
 350 and this can make subcritical cycles more attractive from an economic perspective [62]. Nonetheless, future  
 351 efforts should extend the existing CAMD-ORC model to transcritical cycles.

352 In comparison, when  $C_n = 6$ , the critical temperature of the working fluid is increased, which means the  
 353 the evaporation temperature is no longer constrained by the critical temperature. Instead, the evaporation  
 354 temperature, and therefore evaporation pressure, is constrained by the heat-source temperature profile, and  
 355 the imposed pinch point at the evaporator inlet. This results in no superheating and a lower reduced  
 356 evaporator pressure. Whilst the former means a superheater is no longer required, the latter results in a  
 357 larger latent-heat of evaporation, which impacts the cycle in two ways. Firstly, the larger latent-heat means  
 358 that a larger proportion of the heat-addition process occurs at a constant temperature, which increases the  
 359 average temperature difference between the heat source and working fluid, resulting in more irreversibility.  
 360 Secondly, the larger latent-heat also means that the preheater inlet is no longer pinched, which means this  
 361 cycle absorbs less heat from the heat source. These combined effects result in a 13% reduction in power  
 362 output compared to the optimal cycle.

363 Finally, where  $C_n = 5$ , the maximum power is obtained. This cycle has a high-reduced pressure, minimal  
 364 superheating, and the minimum allowable pinch point is once again observed at both the preheater inlet and

365 the evaporation inlet. Overall, this means that the ORC absorbs the maximum amount of heat possible,  
 366 whilst a low latent heat of vaporisation, and minimal superheating results in the majority of heat-transfer  
 367 occurring during in the preheating region. This minimises irreversibilities within the heat-addition process,  
 368 and results in the maximum power output.

369 The effect of the working fluid, in terms of the number of carbon atoms, on the evaporator and condenser  
 370 thermal load has been reported in Figures 8 and 9 respectively. For the evaporator,  $\dot{Q}_{ph}$ ,  $\dot{Q}_{ev}$  and  $\dot{Q}_{sh}$  refer  
 371 to the preheating, two-phase evaporation, and superheating loads respectively, and for the condenser  $\dot{Q}_{ds}$   
 372 and  $\dot{Q}_{co}$  refer to the desuperheating and two-phase condensation loads respectively. In Figures 8 and 9, only  
 373 the results for one particular working-fluid family have been presented for each heat-source temperature,  
 374 and this corresponds to *n*-alkane family for the 150 °C heat source, and the 2-alkene family for both the  
 375 250 and 350 °C heat sources. However, there was not observed to be a large difference in the breakdown in  
 376 the heat-exchanger load as the working-fluid family is changed, and therefore the discussion in the following  
 377 paragraphs is relevant to all of the families considered.

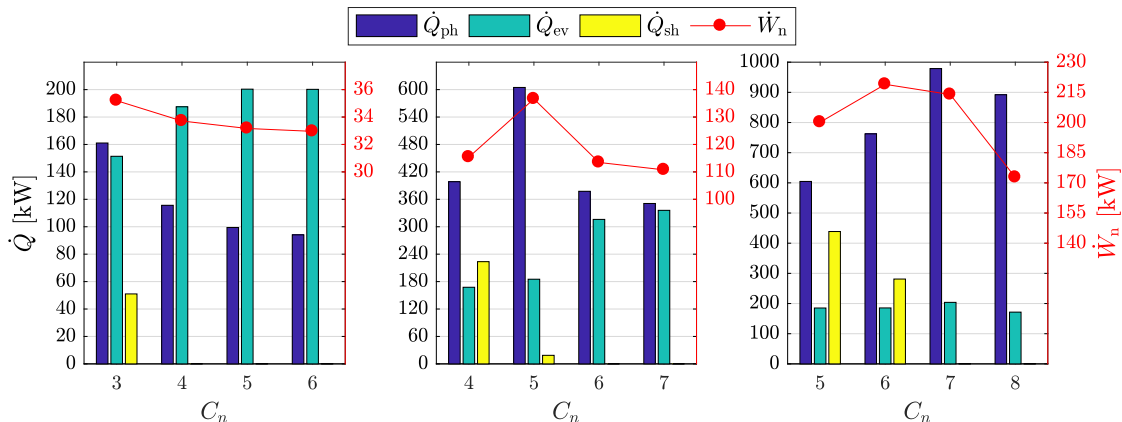


Figure 8: Breakdown of evaporator load for an optimal ORC system operating with different hydrocarbon working fluids. From left to right:  $T_{hi} = 150$  °C (*n*-alkane family);  $T_{hi} = 250$  °C (2-alkene family);  $T_{hi} = 350$  °C (2-alkene family).

378 In terms of the evaporator load a number of observations can be made. Firstly, for the 150 and 250 °C  
 379 heat-source temperatures there is a clear link between maximising the power output and increasing the  
 380 preheater load, with both parameters showing the same trend as  $C_n$  is increased. Moreover, for molecules  
 381 that are less complex than the optimal fluid it is always necessary to have superheating, whilst for molecules  
 382 that are more complex than the optimal fluid the evaporation load increases. Furthermore, it is observed that  
 383 as the heat-source temperature increases the proportion of heat-addition that occurs within the preheater  
 384 increases. More specifically, for the 150 °C heat-source temperature the preheater accounts for between  
 385 32.0% and 44.3% of the total evaporator load, whilst for the 350 °C heat-source temperature, the preheater  
 386 accounts for between 49.2% and 83.8% of the total evaporator load, depending on the fluid.

387 Referring to Figure 9, similar observations for the condenser load can be observed. Firstly, for the 150  
 388 and 250 °C heat-source temperatures, it is observed that maximising the power output also corresponds to

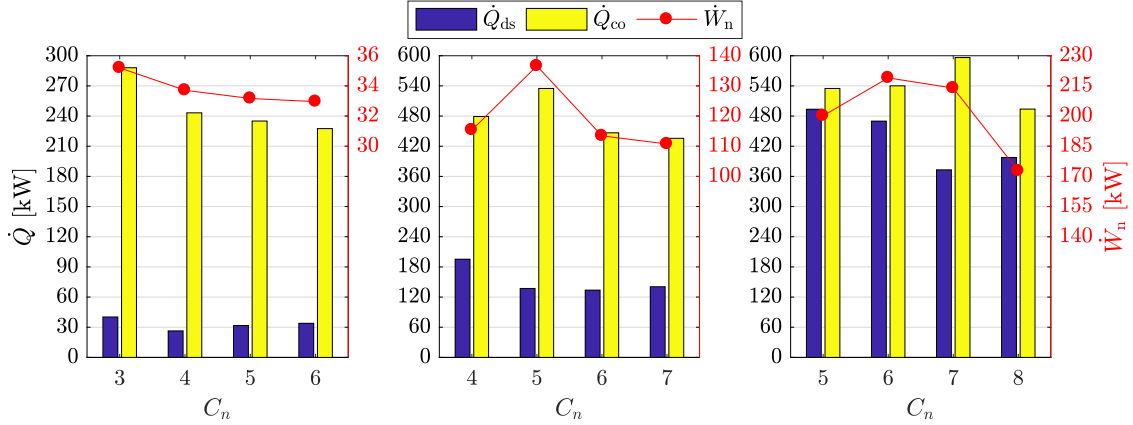


Figure 9: Breakdown of condenser load for an optimal ORC system operating with different hydrocarbon working fluids. From left to right:  $T_{hi} = 150\text{ }^{\circ}\text{C}$  (n-alkane family);  $T_{hi} = 250\text{ }^{\circ}\text{C}$  (2-alkene family);  $T_{hi} = 350\text{ }^{\circ}\text{C}$  (2-alkene family).

389 the largest condensation load, although the difference between the different fluids is not as significant. It  
390 is also observed that in general the less complex molecules result in the highest desuperheater loads. This  
391 was to be expected as these cycles include superheaters, and therefore the working-fluid conditions at the  
392 outlet of the expander will be more superheated than the optimal cycles. In terms of the effect of the  
393 heat-source temperature on the distribution of the condenser load, it is observed that increasing the heat-  
394 source temperature results in a larger proportion of the heat rejection occurring during the desuperheater  
395 stage. For example, for the  $150\text{ }^{\circ}\text{C}$  the desuperheater accounts for between 9.8% and 13.0% of the total  
396 condenser load, whilst for the  $350\text{ }^{\circ}\text{C}$  heat-source temperature this increases to between 38.5% and 48%.  
397 This effect can be explained by considering the behaviour of the saturation dome of hydrocarbon working  
398 fluids as the critical temperature is increased. In general, the saturation dome of a working fluid with a  
399 higher critical temperature will have a larger overhang when viewed on a  $T$ - $s$  diagram. Therefore, expansion  
400 will result in a larger amount of superheat at the expander outlet. Moreover, this effect becomes more  
401 pronounced as the pressure ratio is increased, as is the case as the heat-source temperature increases. The  
402 increased desuperheater load for the  $350\text{ }^{\circ}\text{C}$  heat-source temperature also has an effect on the thermal  
403 efficiency, as increased desuperheating raises the average temperature of heat rejection. This, coupled to  
404 higher condensation temperatures for the  $350\text{ }^{\circ}\text{C}$  systems owing to the fixed heat-sink heat capacity rate,  
405 means that despite the  $350\text{ }^{\circ}\text{C}$  systems producing significantly more power, the thermal efficiencies are  
406 similar to the  $250\text{ }^{\circ}\text{C}$  systems. More specifically, the thermal efficiencies range between 9.7% and 11.2% for  
407 the  $150\text{ }^{\circ}\text{C}$  systems, 14.6% and 16.9% for the  $250\text{ }^{\circ}\text{C}$  systems and 16.2% and 18.1% for the  $350\text{ }^{\circ}\text{C}$  systems.

#### 408 4.3. Component sizing performance

409 Following from the thermodynamic analysis, the required heat-transfer areas for the evaporator and con-  
410 denser can be obtained using the heat-exchanger sizing model based on the group-contribution transport  
411 properties. In Figures 10 and 11 the breakdown of the evaporator and condenser heat-transfer area require-

412 ments are plotted for the same working fluids and cycles considered in Figures 8 and 9. For the evaporator,  
 413  $A_{ph}$ ,  $A_{ev}$  and  $A_{sh}$  refer to the preheating, two-phase evaporation, and superheating areas respectively, and  
 414 for the condenser  $A_{ds}$  and  $A_{co}$  refer to the desuperheating and two-phase condensation areas respectively.

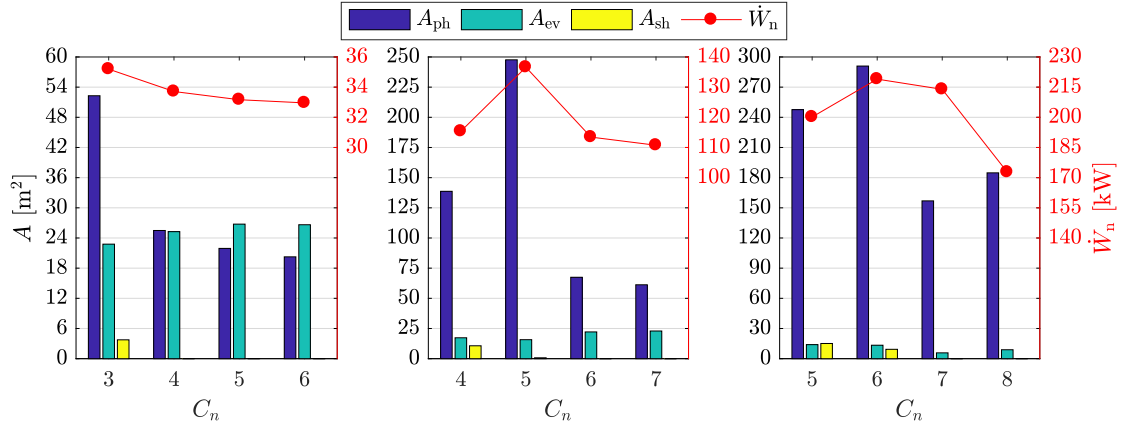


Figure 10: Breakdown of evaporator heat-transfer area requirements for an optimal ORC system operating with different hydrocarbon working fluids. From left to right:  $T_{hi} = 150\text{ }^{\circ}\text{C}$  ( $n$ -alkane family);  $T_{hi} = 250\text{ }^{\circ}\text{C}$  (2-alkene family);  $T_{hi} = 350\text{ }^{\circ}\text{C}$  (2-alkene family).

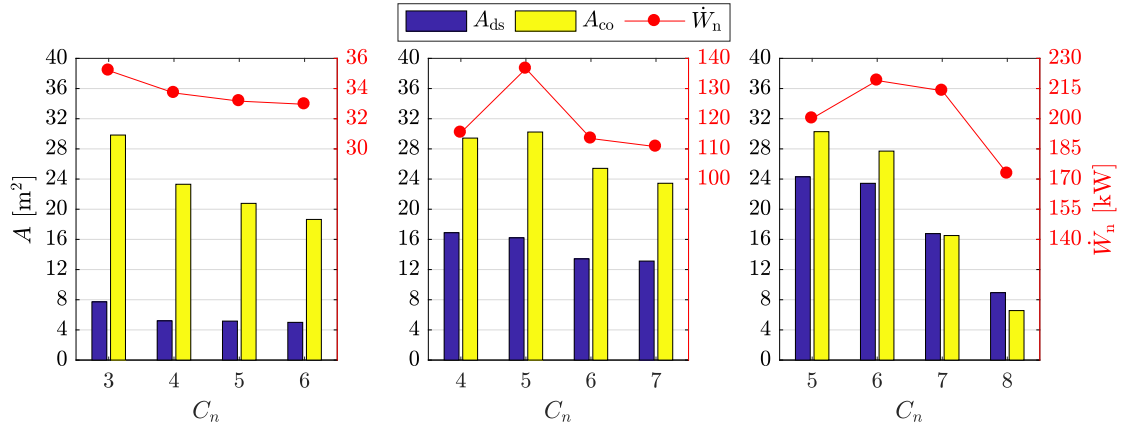


Figure 11: Breakdown of condenser heat-transfer area requirements for an optimal ORC system operating with different hydrocarbon working fluids. From left to right:  $T_{hi} = 150\text{ }^{\circ}\text{C}$  ( $n$ -alkane family);  $T_{hi} = 250\text{ }^{\circ}\text{C}$  (2-alkene family);  $T_{hi} = 350\text{ }^{\circ}\text{C}$  (2-alkene family).

415 Unsurprisingly, for each heat-source temperature, the cycle with the highest power output results in the  
 416 highest heat-transfer area requirements for the evaporator, corresponding to 78.8, 264.1 and 313.6  $\text{m}^2$  for  
 417 the  $n$ -propane, 2-pentene and 2-hexene cases respectively. However, it is observed that whilst selecting a  
 418 different working fluid will cause a reduction in the power output, the reduction in the heat-transfer area  
 419 can be significant. For example, for the three heat-source temperatures, if  $C_n$  is increased by one, the power  
 420 output is reduced by 4.2%, 16.9% and 2.3%, but this corresponds to a reduction in the total evaporator area  
 421 by 35.6%, 66.1% and 48.1% respectively. Therefore, it is clear that a trade-off exists that must be considered

422 when selecting the most suitable working fluid for a particular application.

423 Considering the breakdown of the evaporator heat-transfer area, it is observed that in general the pre-  
424 heater section accounts for the largest percentage of the required area. This was to be expected from con-  
425 sidering the evaporator load breakdown (Figure 8), but is further exaggerated since the overall heat-transfer  
426 coefficient for two-phase evaporation is generally higher than it is for single-phase heat transfer, meaning  
427 a larger area is required to transfer the same amount of heat. For all the fluids evaluated, the preheating  
428 overall heat-transfer coefficient ranged between 176 and 305 W/(m<sup>2</sup> K), whilst the two-phase evaporation  
429 overall heat-transfer coefficient ranged between 268 and 591 W/(m<sup>2</sup> K). Adding to this, the minimum pinch  
430 point is recorded in the preheater section reducing significantly the log-mean temperature difference between  
431 the two working fluids. This results in an increase of the area requirements of the preheater in comparison  
432 to the two-phase evaporating section, even for very similar heat-transfer loads. A case in point is given  
433 by fluids with  $C_n = 3$  at the 150 °C heat-source temperature that have similar preheater and evaporator  
434 loads (Figure 8), but the preheater area required is more than double the respective one for the evaporator  
435 section (Figure 10). Similar findings are observed for fluids with  $C_n = 7$  at 250 °C heat-source temperature.  
436 Referring to the results in Figure 10, for the 150 °C heat-source temperature the preheater accounts for  
437 between 43.2% and 66.4% of the total evaporator area, whilst for the 350 °C heat-source temperature, the  
438 preheater accounts for between 89.5% and 96.4% of the total evaporator area, depending on the fluid.

439 For the condenser heat-transfer area requirements, similar observations to those made when evaluating  
440 the condenser load are found; namely that, with the exception of the  $C_n = 5$ , 350 °C case study, the thermo-  
441 dynamic optimal cycles result in the largest heat exchangers. More specifically, for the *n*-propane, 2-pentene  
442 and 2-hexene cases, the total condenser areas are 37.6, 46.4 and 51.1 m<sup>2</sup> respectively. Interestingly though,  
443 it is observed the required condenser area doesn't increase significantly as the heat-source temperature in-  
444 creases. This is attributed to the higher temperature differences between the heat sink, and the expander  
445 outlet temperature and the condensation temperature as the heat-source temperature increases. For exam-  
446 ple, for the cycles reported in Figure 11, the condensation temperatures range between 303.8 and 306.7 K for  
447 the 150 °C heat source, 313.7 and 318.5 K for the 250 °C heat source and 318.5 and 354.5 K for the 350 °C  
448 heat source. This significant increase in the condensation temperature increases the log-mean temperature  
449 difference, and therefore heat flux, in the condenser resulting in much lower heat-transfer area requirement  
450 for a similar load. It is also worth noting that the significant increase in the condensation temperature  
451 for the  $C_n = 8$ , 350 °C case, is because a minimum condensation pressure constraint is applied (0.25 bar)  
452 during the optimisation. In fact, for both the  $C_n = 7$  and  $C_n = 8$  cases for this heat-source temperature  
453 the condensation pressure is actually equal to the minimum allowable condensation pressure. Therefore, a  
454 lower condensation temperature cannot be achieved without violating this constraint. Not only does this  
455 have a significant effect on the size condenser area, as observed in Figure 11, it also has an impact on the  
456 evaporator area requirements, as the minimum allowable heat-source temperature must also increase, which  
457 in turn moves the evaporator pinch-point to the preheating inlet, rather than at the start of evaporation.



458 Comparing the breakdown of the condenser heat-transfer area requirements, and the breakdown of the  
 459 condenser load, it is observed that the breakdown of the load and area are fairly similar. The desuperheater  
 460 area accounts for a slightly larger proportion of the total condenser area, compared to the desuperheating  
 461 load, and this can again be attributed to the higher overall heat-transfer coefficients for two-phase heat  
 462 transfer compared to the single-phase heat transfer. For all the fluids evaluated, the desuperheating overall  
 463 heat-transfer coefficients ranged between 385 and 518 W/(m<sup>2</sup> K), whilst for two-phase condensation it ranged  
 464 between 926 and 1,450 W/(m<sup>2</sup> K).

465 The total heat-transfer area requirements (*i.e.*, total evaporator area  $A_h$ , and total condenser area  $A_c$ )  
 466 for each heat-source temperature and each working fluid considered within this study are plotted in Figure  
 467 12. Considering this figure, and referring back to Figure 6, it is clear that the optimal thermodynamic cycles  
 468 always result in the largest heat exchangers, and this is particularly true for the 150 and 250 °C heat-source  
 469 temperatures. The reason can also be explained by reconsidering Figure 7, and the accompanying discussion.  
 470 That is to say that the optimal thermodynamic cycle results in a large preheating load, which means a large  
 471 proportion of the available heat is absorbed by the cycle which increases power output. However, this heat  
 472 transfer occurs under a small temperature difference, resulting a large heat-transfer area requirement.

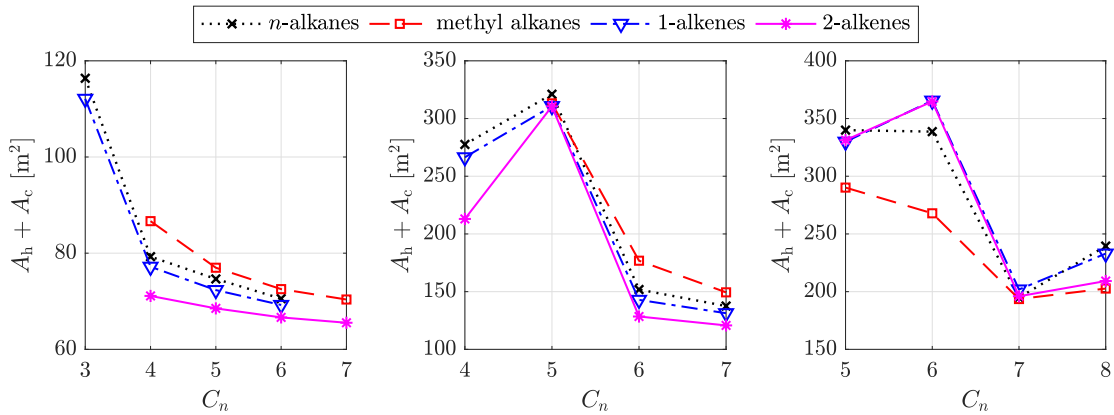


Figure 12: Total heat-transfer area requirements for each cycle previously identified in Figure 6. From left to right:  $T_{hi} = 150$ , 250, 350 °C.

473 Finally, to conclude this section it is useful to evaluate the expander volume ratio, defined as the ratio  
 474 of the inlet and out densities (*i.e.*  $\rho_3/\rho_4$ ). For the 150 and 250 °C systems, the volume ratio increases as  
 475 the number of carbon atoms is increased, but is not found to vary significantly when comparing the different  
 476 fluid families. More specifically, the expansion ratios range between 3.1 to 9.7 for the 150 °C systems, and 6.7  
 477 and 36.1 for the 250 °C systems. For the 350 °C system, the volume ratio increases until the condensation  
 478 pressure constraint comes into play ( $C_n > 7$ ), after which it reduces. For these systems the volume ratio is  
 479 found to range between 17.5 and 162. Ultimately, the volume ratios for the 150 and 250 °C systems can be  
 480 accommodated by a single-stage radial turbine, whilst it is likely that the 350 °C systems, operating with  
 481 working fluids of increasing molecular complexity, would be more suited to a multi-stage design. Therefore,

482 whilst it is reiterated that expander design is not a focus of this paper, future research should account for  
 483 the effect of the volume ratio on both the expander design, and associated cost, in the future.

#### 484 4.4. Thermoeconomic results

485 Clearly, there is a trade-off between thermodynamic performance and the size of the system components.  
 486 Using the known heat-transfer areas, the pump work and expander work for each cycle, the cost correlations  
 487 described in Section 2.4 can be used to obtain the specific-investment cost (SIC) (Figure 13). Within this  
 488 study, the heat-source heat capacity rate has been fixed at 4.2 kW/K and the heat-source temperature  
 489 has been varied, which as observed from Figure 6, has led to different sized systems for each heat-source  
 490 temperature. Therefore, when evaluating the cost of the system there are two factors at play; the size of  
 491 the system, and the heat-source temperature. On the one hand, larger systems will be associated with lower  
 492 relative costs for the manufacturing of components, owing to economy-of-scale effects, which will reduce the  
 493 SIC. On the other hand, higher temperature systems will be associated with higher power outputs, owing  
 494 to higher thermal efficiencies, which will also reduce the SIC. Therefore, as one would expect, it is observed  
 495 in Figure 13 that the lowest temperature and smallest systems (150 °C) correspond to highest SIC whilst  
 496 the highest temperature and largest systems (350 °C) correspond to the lowest SIC. Unfortunately, it is  
 497 difficult to determine what fraction of the reduction in SIC for the 250 and 350 °C systems can be attributed  
 498 to the increase in the system size, and what fraction can be attributed to the increase in the heat-source  
 499 temperature. Future research should attempt to decouple these two effects, for example by scaling the  
 500 heat-source capacity rate such that the power output from each system is the same.

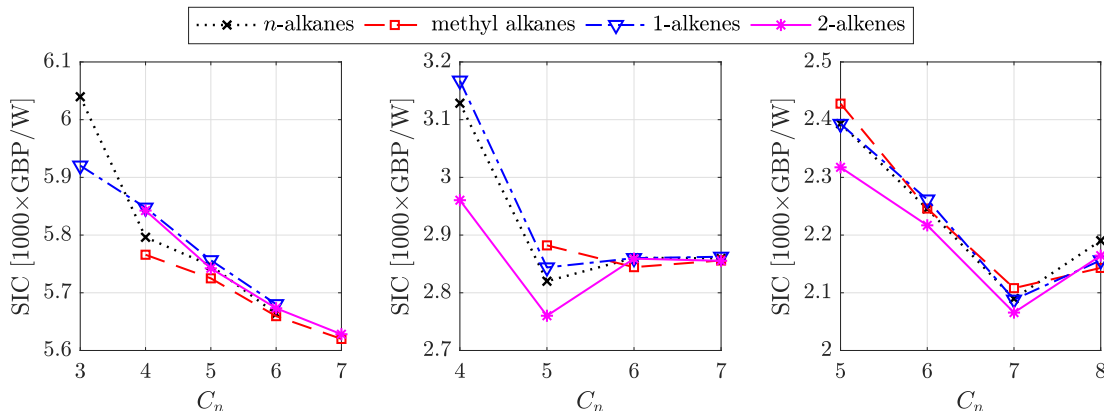


Figure 13: Specific investment cost (SIC) in £/kW for each optimal cycle previously identified in Figure 6. From left to right:  $T_{hi} = 150, 250, 350$  °C.

501 Referring back to Figure 13, it is observed that for each heat-source temperature and hydrocarbon family,  
 502 there appears to be a particular working fluid that will minimise the SIC. For the 150, 250 and 250 °C heat-  
 503 source temperatures the minimum SICs are 5,620, 2,760 and 2,070 £/kW respectively, and these are found  
 504 for  $C_n = 7$  (isoheptane),  $C_n = 5$  (2-pentene) and  $C_n = 7$  (2-heptene) respectively. It should be noted

505 that within the CAMD-ORC model there is no consideration of the order of the functional groups within  
506 the molecule. Therefore, isoheptane refers to either 2-methyl hexane or 3-methyl hexane, depending on the  
507 location of the  $-CH$  group. For the 250 °C heat source 2-pentene is found to both maximise the power output  
508 and minimise the SIC, and is therefore identified as the optimal working fluid. However, for the other two  
509 heat sources, different working fluids are identified based on whether a thermodynamic or technoeconomic  
510 performance metric is used.

511 Of particular interest, are the results for the 350 °C heat source, which suggest that in terms of minimising  
512 the SIC, it could be beneficial to use a working fluid with  $C_n = 7$ . The four fluids considered with  $C_n = 7$   
513 have condensation temperatures ranging between 49.4 °C (methyl alkane) and 56.8 °C (2-alkene), with  
514 corresponding pinch points at the start of evaporation of 53.1 °C and 39.5 °C respectively. This results  
515 in relatively large temperature differences within the heat exchangers, thus reducing the heat-transfer area  
516 requirement and therefore cost. Lowering  $C_n$  reduces both the condensation temperature and the pinch  
517 point, resulting in better performance but much higher costs. On the other hand, increasing  $C_n$  to 8  
518 corresponds to condensation temperatures between 74.5 °C (methyl alkane) and 81.7 °C ( $n$ -alkane) resulting  
519 in a significant reduction in performance. It is also interesting to note that there is only a small difference  
520 between the optimal SIC for each fluid family, with the optimal SIC ranging between 2,065 £/kW (2-alkene)  
521 and 2,108 £/kW (methyl alkane), which corresponds to a 2.1% increase in the SIC when using a methyl  
522 alkane compared to a 2-alkene. Ultimately, this suggests that in this case the molecular complexity (*i.e.* the  
523 number of carbon atoms) is more critical than the specific molecular structure.

524 In order to confirm whether the SIC values obtained within this study are representative of actual  
525 ORC systems, the results from this study are compared to SIC data available within the literature. More  
526 specifically, Lemmens [53] collated cost data for ORC systems designed for different applications, including  
527 biomass, solar geothermal and waste-heat recovery. In Figure 14 the results from the current study are  
528 compared to the data reported by Lemmens for waste-heat recovery applications, adjusted from €<sub>2014</sub> to  
529 £<sub>2017</sub> using the CEPCI values for 2014 (576.1) and 2017 (562.1), and the current exchange rate (€ 1 = £ 0.87).

530 From Figure 14 it is observed that the SIC values obtained within this paper match well with those  
531 reported within the literature, and this is particularly true for the 250 and 350 °C systems. The SIC values  
532 obtained for the 150 °C systems follow the general trend in that SIC increases as the system size reduces, but  
533 are slightly higher than SIC values taken from the literature. However, it is worth noting that Lemmens did  
534 not consider the effect of heat-source temperature on the system economics. In reality, a low-temperature  
535 heat source will lead to a lower thermal efficiency, and therefore it is reasonable to assume a 50 kW, 150 °C  
536 system will have a higher SIC than a 50 kW, 250 °C system. Therefore, the heat-source temperature is  
537 actually a third dimension, which is not reported Figure 14. With this in mind, it is reasonable to accept  
538 the SIC values obtained for the 150 °C systems.

539 Alongside considering the SIC values obtained for the systems, it is also interesting to consider the  
540 breakdown in the system cost, and this is reported in Figure 15 for the same working fluids previously

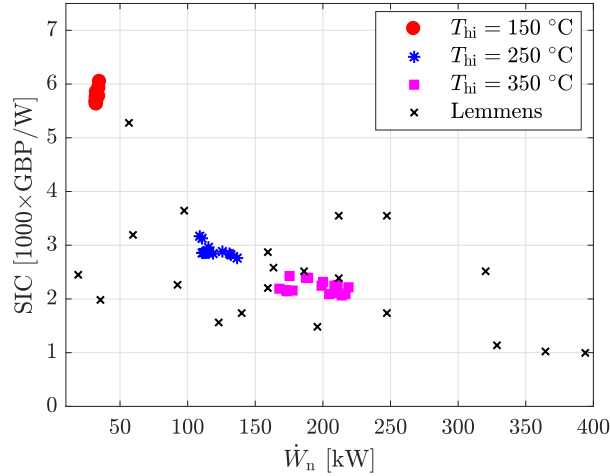


Figure 14: Effect of the system size (in kW) on the specific-investment cost in  $\text{£}/\text{kW}$ . The results from this study for the three different heat-source temperatures are compared to SIC data reported by Lemmens [53].

541 evaluated in terms of the heat-exchanger load and heat-transfer area requirements for the evaporator and  
 542 condenser. Again, it is noted that the results reported in this figure are representative of the results obtained  
 543 for each working-fluid family.

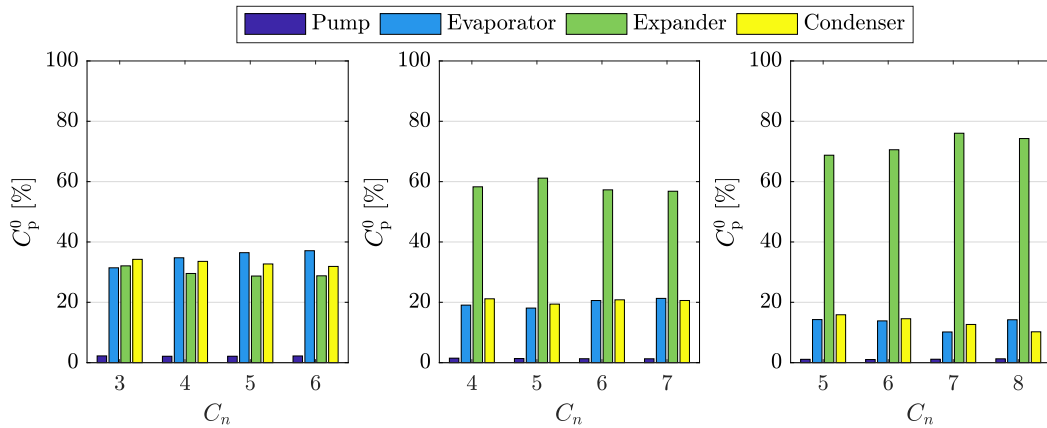


Figure 15: Breakdown of the system cost system for different hydrocarbon working fluids and heat-source temperatures. From left to right:  $T_{hi} = 150^\circ\text{C}$  ( $n$ -alkane family);  $T_{hi} = 250^\circ\text{C}$  (2-alkene family);  $T_{hi} = 350^\circ\text{C}$  (2-alkene family).

544 Firstly, it is noted that for all the cases considered the pump only accounts for a very small percentage  
 545 ( $< 2.3\%$ ) of the total system cost. Furthermore, the evaporator and condenser both account for a similar  
 546 percentage of the overall costs, corresponding to approximately 35%, 20% and 13% for the 150, 250 and  
 547 350  $^\circ\text{C}$  heat-source temperatures respectively. However, the most obvious observation from Figure 15 is  
 548 the significant percentage of the total cost that the expander accounts for as the heat-source temperature  
 549 increases. This behaviour can, in part, be explained since a higher heat-source temperature will lead to  
 550 a higher cycle efficiency. Therefore, a greater percentage of the heat that is input into the system can

551 be converted into power, which therefore requires an expander with a higher power output, and a larger  
552 generator, for the same rating of heat exchanger.

553 Another possible reason for such a large cost estimate for the expander could relate to the suitability of  
554 the expander cost correlation for ORC systems. Arguably, within an ORC system, the cost of the expander  
555 is the largest unknown, particular for small-scale systems below a few-hundred kW, as the commercialisation  
556 of these systems is still in its infancy. For this study, the material factor  $F$  for the expander, which accounts  
557 for component manufacturing, was set to 3.5 based on recommendations within the literature. However, it  
558 should be noted that these correlations were not developed specifically for ORC expanders but they originate  
559 from the chemical industry, and should be used for comparing alternative system configurations and working  
560 fluids performance, where the relative results are more important than the absolute cost figures.

561 Despite possible uncertainties with the cost correlations, referring back to Figure 14 it has been shown  
562 that the SIC values predicted by the CAMD-ORC are in good agreement with values reported within the  
563 literature. Moreover, it should be stated the primary aim of this paper has been to develop a CAMD-  
564 ORC framework that can be used to identify novel ORC architectures from a thermoeconomic perspective.  
565 Therefore, the cost correlations applied within this framework can be easily adapted as the ORC market  
566 continues to grow, and more cost information on the system components becomes available.

## 567 **5. Further economic perspectives**

568 In this paper, we assume a constant heat-source mass-flow rate. However, in several applications this heat  
569 stream is variable both in mass-flow rate and temperature level, on the basis of the specific production  
570 process (*i.e.* intermittent waste heat from food and other industrial processes, variable solar energy, seasonal  
571 biomass supply and so on). Moreover, this heat stream could be used not only to produce electricity using  
572 an ORC, but also to match on-site heating/cooling demand at different temperature levels, according to the  
573 typology of energy demand (*i.e.* commercial/residential or industrial). In some cases, this means that the  
574 ORC system configuration should be optimised to maximise the global energy-conversion efficiency, instead  
575 of the electrical efficiency. This could include using the discharged heat from the ORC cooling stream for  
576 further on-site cogeneration, or accounting for smart operating strategies to modulate or switch on/off the  
577 ORC in order to follow the heat demand. In addition, the intermittency of heat source introduces further  
578 trade-offs in the optimal thermal-storage capacity, considering that thermal storage could increase the ORC  
579 operating hours, but increases costs and reduces the ORC input temperature and, in turn, the ORC electric  
580 conversion efficiency.

581 The overall thermodynamic performance of the ORC should be optimised for different outlet temperatures  
582 of the ORC cooling stream. A higher outlet temperature increases the energy of the heat-sink stream but  
583 decreases the power output of the expander. Conversely, a low outlet temperature allows for a high power  
584 output, but has a low potential to heat buildings or match other industrial thermal energy demand. Moreover,  
585 the waste-heat supply and the low temperature heating demand profiles are often not well matched. This

586 means that, without a proper thermal storage system, cogenerated heat from the ORC-CHP can be wasted  
587 over large periods of the year. This is particularly true when waste heat availability makes profitable a  
588 base-load CHP operation, instead of thermal load following operations. The optimal working fluid for a  
589 given temperature of heat demand identified from an optimisation procedure may not be the optimal one  
590 if the heating demand is affected by high temporal variations, and does not match the CHP output profile.  
591 For this reason, the influence of heat-demand profile on optimal working fluid selection and global CHP  
592 conversion efficiency should be taken into account.

593 Some of these trade-offs have been addressed in recent literature, such as in Ref. [63], which includes  
594 the optimisation of the ORC working fluid as the temperature of heat demand and the operational model  
595 change. Other studies compare leveled costs of energy and profitability of ORC configurations as a function  
596 of intermittency of heat source [64], and thermal-energy storage size and intermittency of solar energy input  
597 [65]. Operational strategies and the dynamics of heat supply-energy demand have also been studied [66],  
598 in addition to the possibility to match electric, heating and cooling demand via smart operation of ORC  
599 coupled to heat sources at different temperature levels [67].

600 In light of these considerations, the next step for waste heat ORC applications and working-fluid op-  
601 timisation should consider the system operational strategies (*i.e.* minimising the levelised-cost of energy  
602 instead of the SIC), the ORC cooling stream temperature, the matching of heat discharged from the ORC  
603 to on-site heat demand, and the dynamics between heat sources and energy demand. Moreover, the effect of  
604 the condensing temperature on the condenser size, and in turn, the specific-investment cost needs further in-  
605 vestigation to understand the sensitivity of the system to this parameter. Finally, studies should address the  
606 broader benefits in terms of energy systems flexibility that could be provided by such distributed waste-heat  
607 recovery options.

## 608 **6. Conclusions**

609 The discovery of new working-fluids that can improve performance while meeting increasingly restrictive  
610 environmental legislation, and the identification of novel and optimal ORC systems based on technoeconomic  
611 performance indicators are key steps to enable a more widespread uptake of ORC technology. The aim of  
612 this paper has been to incorporate technoeconomic analysis, through component sizing and suitable cost  
613 correlations, into an existing CAMD-ORC framework, based on the SAFT- $\gamma$  Mie equation of state.

614 Discretised heat-exchanger sizing models, based on group-contribution methods for determining transport  
615 properties, have been developed to size the evaporator and condenser for optimised ORC systems, and the  
616 resulting specific-investment costs (SIC) have been determined using suitable cost correlations. In addition,  
617 the existing CAMD-ORC framework has been extended to allow the consideration of novel cycle architectures,  
618 including recuperated and partially-evaporated cycles, and cycles operating with working-fluid mixtures.  
619 Both the thermodynamic model and heat-exchanger sizing models have been validated against data from  
620 NIST REFPROP, and a good agreement is found for the working-fluids considered. The largest deviations

621 observed in the heat-exchanger area were +18% and -13%, when sizing the condenser for 2-butene, and  
622 the evaporator for propene respectively. These relatively small deviations confirm the suitability of the  
623 group-contribution transport property prediction methods.

624 From the case study, it is found that working fluids that maximise the power output from the system  
625 generally have the highest heat-exchanger area requirements. Therefore, working-fluid selection based on  
626 SIC minimisation can result in different optimal working fluids to those identified from an optimisation that  
627 considers power output or other common thermodynamic objective functions. For the three heat-source  
628 temperatures considered (150, 250 and 350 °C, each with  $\dot{m}c_p = 4.2$  kW/K corresponding to a small to  
629 medium-scale application) the three working fluids that minimise the SIC are isoheptane, 2-pentene and  
630 2-heptene, with SICs of 5,620, 2,760 and 2,070 £/kW respectively. The corresponding power outputs for  
631 these systems are 32.9, 136.6 and 213.9 kW, and these power outputs are 6.38%, 0.0% and 2.32% lower  
632 than the power outputs obtained for working fluids that maximise the power output. This corresponds to  
633 a reduction in the SIC of 6.95%, 0.0% and 6.82%. Overall, these results demonstrate the importance of  
634 considering technoeconomic performance within the CAMD-ORC framework, and three optimal working  
635 fluids have been identified for different heat-source temperatures.

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845 **Nomenclature**

846 **Abbreviations**

847 CAMD Computer-aided molecular design

848 ORC Organic Rankine cycle

849 SAFT Statistical associating fluid theory

850 SIC Specific-investment cost, £/kW

851 **Greek Symbols**

852  $\epsilon_r$  Recuperator effectiveness

853  $\eta$  Isentropic efficiency

854  $\mu$  Dynamic viscosity, Pa s

855  $\rho$  Density, kg/m<sup>3</sup>

856 **Roman Symbols**

857  $\Delta T_{\log}$  Counter-flow log-mean temperature difference, K

858  $\Delta T_{\text{sh}}$  Degree of superheating, K

859  $\dot{m}$  Mass flow rate, kg/s

860  $\dot{Q}$  Heat exchanger load, J/s

861  $\dot{W}$  Power, J/s

862  $PP$  Pinch point, K

863  $Re$  Reynolds number

864  $A$  Heat-transfer area, m<sup>2</sup>

865  $C_p^0$  Component cost, £

866  $C_n$  Number of carbon atoms

867  $c_p$  Specific heat capacity at constant pressure, J/(kg K)

868  $D$  Diameter, m

869  $F$  Material factor

870	$f$	Friction factor
871	$h$	Enthalpy, J/kg
872	$k$	Thermal conductivity, W/(m K)
873	$L$	Length, m
874	$p$	Pressure, Pa
875	$p_r$	Reduced pressure
876	$s$	Entropy, J/(kg K)
877	$T$	Temperature, K
878	$U$	Overall heat-transfer coefficient, W/(m <sup>2</sup> K)
879	$u$	Velocity, m/s
880	$x$	Mass-fraction of fluid 1 in a two-fluid mixture
881	$Z$	Cost coefficient
882	$z$	Expander inlet design parameter

883 **Subscripts**

884	1-4	ORC state points
885	c	Heat sink/condenser
886	co	Condensation
887	cr	Critical point
888	ds	Desuperheating
889	e	Expander
890	ev	Evaporation
891	h	Heat source/evaporator
892	n	Net
893	o	Working fluid
894	p	Pump
895	ph	Preheating
896	sh	Superheating

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