Cell Formation Problem Solved Exactly with the Dinkelbach Algorithm

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Abstract. The cell formation problem has been extensively studied in the literature, but very few authors have proposed exact methods to solve it. In this paper the cell formation problem is transformed into an equivalent binary linear fractional programming problem, and it is solved exactly with the Dinkelbach algorithm. At each iteration of the method, the binary linear problem is solved with the IBM ILOG CPLEX 12.2 Optimizer. A numerical experimentation is completed using 35 benchmark problems, and 27 of these problems are solved exactly. Moreover, these results allow us to validate the quality of the solution generated with metaheuristic methods proposed in the literature since the best known solutions are equal to the optimal solutions for these 27 problems. To our knowledge, this is the first time that experimentation with an exact procedure is completed for the 35 benchmarked problems used in the literature.

Keywords. Cell formation problem, binary linear fractional programming, grouping efficiency, Dinkelbach algorithm, optimal solution

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1 Introduction

In group technology or in cellular manufacturing, a system including machines and parts are interacting. To maximize the efficiency of the system, a cell formation problem is solved in order to partition the system into subsystems that are as autonomous as possible in the sense that the interactions of the machines and the parts within a subsystem are maximized, and that the interactions between machines and parts of other subsystems are reduced as much as possible. This gives rise to solving a cell formation problem.

The cell formation problem is a NP hard optimization problem [1]. For this reason, several metaheuristic methods have been developed over the last forty years to generate good solutions in reasonable computational time. Nevertheless, only few integer programming formulations [2, 3, 4] have been proposed to solve the smaller problems in order to evaluate the quality of the solutions proposed by the metaheuristic methods. In this paper the cell formation problem is transformed into an equivalent binary linear fractional programming problem, and it is solved exactly with the Dinkelbach algorithm. At each iteration of the method, the binary linear problem is solved with the IBM ILOG CPLEX 12.2 Optimizer. Furthermore, we can evaluate how close the solutions generated by metaheuristic methods are from optimality.

In order to learn more about the different methods proposed to solve the cell formation problem, we refer the reader to the survey proposed in [5], where the authors review briefly the different methodologies used to solved the problem: cluster analysis, graph partitioning, mathematical programming, genetic (population based) algorithms, local search methods (tabu search, simulated annealing), and hybrids of these methods. More recently two additional review papers of the cell formation problem have been published. In the first paper Papaioannou and Wilson [6], the authors survey the different techniques used to solve the problem, and they classify the different solution techniques as follows:

- Cluster analysis: techniques for recognizing structure in a data set
- Graph partitioning approaches: where a graph or a network representation is used to formulate the cell formation problem
- Mathematical programming methods: the cell formation problem is formulated like a non linear or linear integer programming problem
Heuristic, Meta-heuristic and hybrid Meta-heuristic: The most popular methods are: simulated annealing, tabu search, genetic algorithms, colony optimization, particle swarm optimization, neural networks and fuzzy theory.

In the second paper Ghosh et al. [7], the authors introduce a survey of various genetic algorithms used to solve the cell formation problem. The success of genetic algorithms in solving this problem induced researchers to consider different variants and hybrids in order to generate very robust techniques.

To determine the best-known solutions for the 35 benchmark problems, we consider the results reported in Goncalves and Resende [5], James et al. [8], Tunnukij and Hicks [9], Mahdavi et al. [10], Nokhtehdan et al. [11], Xiangyong et al. [12] where the authors compare their results with other already published in the literature. Hence for each problem we consider the best-known solution among those generated by the following methods:

- GRAPHICS [clustering method] Srinivasan and Narendran [14]
- GATSP [genetic algorithm solving a traveling salesman formulation] Cheng et al. [16]
- SA [simulated annealing] Zolfaghari and Liang [18], Lin et al. [19], Ying et al. [20]
- GA [genetic algorithm] Zolfaghari and Liang [18]
- TS [tabu search] Zolfaghari and Liang [18]
- HGGA [hybrid grouping genetic algorithm] James et al. [8]
- EnGGA [grouping genetic algorithm and greedy heuristic] Tunnukij and Hicks [9]
- ACO-CF [ant colony optimization metaheuristic] Xiangyong et al. [12], Farahani and Hosseini [21].

To complete this analysis, we consider a set of 35 benchmark problems that the authors solve to compare their results. In a previous study, the authors in [22] solve a linear approximation of the cell formation problem using the CPLEX Optimizer reaching an optimal solution of their approximation problem in 22 of the 35 benchmarked problems. In this paper, we introduce an exact method to solve the cell formation problem based on the Dinkelbach algorithm [23]. The numerical experimentation indicates the optimal solution is obtained for 27 problems.
In Section 2, we describe the cell formation problem using the grouping efficacy measure used in literature, and we introduce an equivalent formulation as a binary linear fractional programming problem. The Dinkelbach algorithm is described in section 3. The numerical results are summarized in Section 4.

**2 Problem formulation**

To formulate the cell formation problem, consider the following notation:

- \( M \): number of production machines.
- \( P \): number of production parts.
- \( C \): number of production cells.
- \( I \): set of \( M \) machines \((i = 1, \ldots, M)\),
- \( J \): set of \( P \) parts \((j = 1, \ldots, P)\),
- \( K \): set of \( C \) cells \((k = 1, \ldots, C)\),

The production incidence matrix \( A = \begin{bmatrix} a_{ij} \end{bmatrix} \) indicates the interactions between the machines and the parts:

\[
a_{ij} = \begin{cases} 1 & \text{if machine } i \text{ process part } j \\ 0 & \text{otherwise.} \end{cases}
\]

Furthermore, a part \( j \) may be processed by several machines. A production cell \( k \ (k = 1, \ldots, C) \) includes a subset (group) of machines \( I_k \subset I \) and a subset (family) of parts \( J_k \subset J \). The problem is to determine a solution including \( C \) production cells \( (I, J) = \{(I_1, J_1), \ldots, (I_C, J_C)\} \) as autonomous as possible. Note that the \( C \) production cells induces partitions of the machines set and of the parts set:

\[
I_1 \cup \ldots \cup I_C = I \quad \text{and} \quad J_1 \cup \ldots \cup J_C = J
\]

and for all pairs of different cell indices \( k_1 \) and \( k_2 \in \{1, \ldots, C\} \)

\[
I_{k_1} \cap I_{k_2} = \emptyset \quad \text{and} \quad J_{k_1} \cap J_{k_2} = \emptyset.
\]

To illustrate the production cells concept, consider a machine-part incidence matrix in Table 1. Table 2 indicates a partition into 3 different cells illustrated in the gray zones. The solution includes the 3 machine groups \{1,4,6\}, \{3,5\}, \{(2)\} and the 3 part families \{(2,4,6,8), (1,7), (3,5)\}. 

---

Table 1: Machine-part Incidence Matrix

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<th>J</th>
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<tbody>
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Table 2: Cell Formation

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<th>Cell 3</th>
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Table 1. Incidence matrix

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<th>5</th>
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</table>

Table 2. Matrix solution

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<tbody>
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<td>0</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>

The exceptional elements (1,5), (6,1), (6,7), (3,3), (5,8) and (2,1) correspond to entries having a value 1 that lay outside of the gray diagonal blocks.

To measure the autonomy of a solution, different measures have been proposed, Sarker and Khan carry out a comparative study in [24]. In this paper we consider the grouping efficacy (Eff) that is usually used by the authors to compare the efficiency of their methods (see [25]):

\[
Eff = \frac{a - a_{Out}}{a + a_{In}} = 1 - \left( \frac{a_{In} + a_{Out}}{a + a_{In}} \right)
\]  

(1)

where \(a = \sum_{i=1}^{M} \sum_{j=1}^{P} a_{ij}\) denotes the total number of entries equal to 1 in the matrix A, \(a_{Out}\) denotes the number of exceptional elements, and \(a_{In}\) is the number of zero entries in the gray diagonal blocks.

The objective function of maximizing \(Eff\) is then equivalent to minimize \(ComEff = \left( \frac{a_{In} + a_{Out}}{a + a_{In}} \right)\).

To specify a mathematical programming formulation for the cell formation problem, we introduce the following binary variables:
for each pair $i = 1, \ldots, M; k = 1, \ldots, C$

$$x_{ik} = \begin{cases} 1 & \text{if machine } i \text{ belongs to cell } k \\ 0 & \text{otherwise} \end{cases}$$

for each pair $j = 1, \ldots, P; k = 1, \ldots, C$

$$y_{jk} = \begin{cases} 1 & \text{if part } j \text{ belongs to cell } k \\ 0 & \text{otherwise} \end{cases}.$$

It is easy to verify that

$$a_{in}^o = a - \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P a_{ij} x_{ik} y_{jk}$$

$$a_{0}^{out} = \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P (1 - a_{ij}) x_{ik} y_{jk},$$

and it follows that the objective function $ComEff$ is a fractional function in $x$ and $y$.

Denote the numerator and the denominator of $ComEff$ as

$$N(x, y) = a_{0}^{out} + a_{i}^{out} = \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P (1 - a_{ij}) x_{ik} y_{jk} + a - \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P a_{ij} x_{ik} y_{jk}$$

$$= a + \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P (1 - 2a_{ij}) x_{ik} y_{jk},$$

and

$$D(x, y) = a + a_{0}^{in} = a + \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P (1 - a_{ij}) x_{ik} y_{jk},$$

respectively.

Hence the objective function can be written as:

$$ComEff(x, y) = \frac{N(x, y)}{D(x, y)} = \frac{a + \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P (1 - 2a_{ij}) x_{ik} y_{jk}}{a + \sum_{k=1}^C \sum_{i=1}^M \sum_{j=1}^P (1 - a_{ij}) x_{ik} y_{jk}}$$ (2)
Then the cell formation problem can be formulated as in the following $M(x, y)$ model:

$$\begin{align*}
\text{Min } \text{ComEff} &= \frac{N(x, y)}{D(x, y)} = a + \sum_{k=1}^{C} \sum_{i=1}^{M} \sum_{j=1}^{P} (1-2a_{ij})x_{ik}y_{jk} \\
&\quad - \sum_{k=1}^{C} \sum_{i=1}^{M} \sum_{j=1}^{P} (1-a_{ij})x_{ik}y_{jk} \\
\text{Subject to } &\quad \sum_{k=1}^{C} x_{ik} = 1 \quad i = 1, \ldots, M \\
&\quad \sum_{k=1}^{C} y_{jk} = 1 \quad j = 1, \ldots, P \\
&\quad \sum_{i=1}^{M} x_{ik} \geq 1 \quad k = 1, \ldots, C \\
&\quad \sum_{j=1}^{P} y_{jk} \geq 1 \quad k = 1, \ldots, C \\
&\quad x_{ik} = 0 \text{ or } 1 \quad i = 1, \ldots, M; k = 1, \ldots, C \\
&\quad y_{jk} = 0 \text{ or } 1 \quad j = 1, \ldots, P; k = 1, \ldots, C
\end{align*}$$

The constraints (3) and (4) ensure that each machine and each part is assigned to exactly one cell, respectively. The constraints (5) and (6) ensure that each cell includes at least one machine and one part (i.e., no empty cell). Finally, the variables are binary in (7) and (8). In our numerical experimentation we fix the number $C$ of cells for each problem to its value in the best-known solution.

Now we use a similar approach as in Mahdavi et al. [4] to transform the problem $M(x, y)$ into a linear binary fractional problem using additional binary variables $w_{ij}^k$ to replace the product $x_{ik}y_{jk}$:

$$\begin{align*}
w_{ij}^k &= x_{ik}y_{jk} \quad i = 1, \ldots, M; j = 1, \ldots, P; k = 1, \ldots, C \\
w_{ij}^k &= 0 \text{ or } 1 \quad i = 1, \ldots, M; j = 1, \ldots, P; k = 1, \ldots, C
\end{align*}$$

To complete the linear formulation of the cell formation problem, we use the following linear relations equivalent to the quadratic relations (9):

$$\begin{align*}
w_{ij}^k - x_{ik} &\leq 0 \quad i = 1, \ldots, M; j = 1, \ldots, P; k = 1, \ldots, C \\
w_{ij}^k - y_{jk} &\leq 0 \quad i = 1, \ldots, M; j = 1, \ldots, P; k = 1, \ldots, C \\
-w_{ij}^k + x_{ik} + y_{jk} &\leq 1 \quad i = 1, \ldots, M; j = 1, \ldots, P; k = 1, \ldots, C
\end{align*}$$

The binary fractional programming problem $M(x, y, w)$ is obtained by including the additional constraints (10), (11), (12), (13) into problem $M(x, y)$. 

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We can now rely on the Dinkelbach [23] approach to solve the linear binary fractional problem $M(x, y, w)$.

### 3 Dinkelbach procedure

To simplify the notation, denote

$$(X,Y,W) = \{(x, y, w) : \text{solutions satisfying contraints (3) – (8), (10) – (13)}\}$$

The Dinkelbach procedure to solve $M(x, y, w)$ requires solving a sequence of problems where the objective function becomes binary linear by including the denominator $D(x, y, w)$ into the numerator $N(x, y, w)$:

$$M(\lambda, x, y, w) = \min_{(x, y, w) \in (X,Y,W)} \frac{N(x, y, w) - \lambda D(x, y, w)}{a + \sum_{k=1}^{M} \sum_{j=1}^{P} (1-a_{ij}) w_{k}^j}$$

for different values of $\lambda$. This linear binary problem is solved with CPLEX 12.2 Optimizer problem.

The Dinkelbach procedure can now be summarized as follows:
**Dinkelbach procedure**

- **Initialization.**
  - Start with an initial solution \((x^0, y^0, w^0) \in (X, Y, W)\)
  - Take \(\lambda_0 := \frac{N(x^0, y^0, w^0)}{D(x^0, y^0, w^0)}\), and \(\zeta := 1\).

- **Step**
  - Solve the problem \(M(\lambda_{\zeta-1}, x, y, w)\)
  - Let \((x^\zeta, y^\zeta, w^\zeta)\) be an optimal solution of this problem.
  - Let \(F(\lambda_{\zeta})\) be the optimal of this problem.
  - **Stopping rule.** If \(F(\lambda_{\zeta}) < \delta\) (optimality tolerance), then STOP: \((x^\zeta, y^\zeta, w^\zeta)\) is an optimal solution of \(M(x, y, w)\) and \(Eff(x^\zeta, y^\zeta)\) is the maximal grouping efficacy.
  - Otherwise, let \(\lambda_{\zeta} := \frac{N(x^\zeta, y^\zeta, w^\zeta)}{D(x^\zeta, y^\zeta, w^\zeta)}\)
  - Let \(\zeta := \zeta + 1\), and go back to **Step**

Since we expect the best-known efficacy value to be close to the optimal value, then we use this value to initialize \(\lambda^0\) in order to reduce the number of iterations of the algorithm. Thus for each problem \(\lambda^0 = \left(1 - \frac{\text{best-known solution}}{100}\right)\).

Finally, the algorithm converges since the sequence \(\{\lambda_{\zeta}\}\) is strictly decreasing (Crouzeix et al.[26]).

**4 Numerical results**

In this paper we consider the 35 benchmark problems that are usually used by authors to compare the efficiency of their solution approaches. Referring to Goncalves and Resende [5], James et al. [8], Tummukij and Hicks [9], Mahdavi et al. [10], Nokhtehdan et al. [11], Xiangyong et al. [12], and Elbenani et al. [27] papers summarizing the results obtained by several authors, we can identify the best-known solution for each of the 35 problems. Our purpose is to solve the models \(M(x, y, w)\) for the cell formation problem and the sub-problems \(M(\lambda, x, y, w)\) with the Dinkelbach algorithm and the CPLEX 12.2 Optimizer, respectively. The optimal solutions obtained for the grouping efficacy \(Eff\) are compared with the grouping efficiency of the best-known solutions of 35 benchmark problems.
The problems are summarized in Table 3. The problem number and the reference where it is specified are included in the first two columns of the table. The size of the problems (values of $M$ and $P$) and the number of cells $C$ are in columns 3, 4, and 5. The best-known value of the grouping efficacy for each of the 35 problems is included in column 6 of the table.

The computational tests are performed on a AMD processor running at 2.2 GHz with 4096 Kilobytes of central memory and 1024 Kilobytes of memory cache. The relative optimality tolerance $\delta$ in the Dinkelbach algorithm is set at the value $10^{-6}$. The grouping efficacy $Eff$ obtained with the Dinkelbach algorithm and the solution time are reported in columns 7 and 8 of the Table 3, respectively.

The results in the Table 3 indicate that the first 11 problems, and the problems P20 and P22 are easy to solve since they requires less than 32 seconds of computational time. The solution approach can also reach the optimal solution for 14 of the larger problems (P12 to P17, P19, P21, P23, P24, P28, P30, P34, and P35), but the solution time can become very large. Unfortunately, it was not possible to solve the 8 other problems (P18, P25 to P27, P29, and P31 to P33) because the memory limit is reached before getting the optimal solutions. For these problems we place a * in the last column of the Table 3, and in column 7, we indicate in italic the value of $Eff$ reached by our method before stopping.

<table>
<thead>
<tr>
<th>Problem number</th>
<th>Problem source</th>
<th>$M$</th>
<th>$P$</th>
<th>$C$</th>
<th>Best-known solution</th>
<th>Dinkelbach $Eff$</th>
<th>Solution time (sec)</th>
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<tr>
<td>P1</td>
<td>King and Nakornchai [28]</td>
<td>5</td>
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<td>593.2</td>
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<tr>
<td>P14</td>
<td>McCormick [41]</td>
<td>16</td>
<td>24</td>
<td>8</td>
<td>53.26</td>
<td>53.26</td>
<td>15130.5</td>
</tr>
</tbody>
</table>
The approach in [22] allows reaching the optimal solution of 22 problems (P1 to P15, P17, P19, P20, P22 to P24, and P35) even if it is solving exactly a binary linear approximation of the cell formation problem with an earlier version of CPLEX. For the additional 5 problems solved exactly with the Dinkelbach methods (P16, P21, P28, P30 and P34), the optimal value is indicated in bold in Table 3. Finally, note that for the 8 problems (P18, P25 to P27, P29, and P31 to P33) where the method stops because of memory limit, the average value of their best known solutions grouping is better by a factor of 0.78 % better over the average corresponding values of $Eff$ reached by our method.

It follows that the metaheuristic methods reaching the best known solutions seem to be very efficient since they can reach the optimal value $Eff$ for the 27 problems solved to optimality by the Dinkelbach algorithm. But nevertheless, even if the Dinkelbach algorithm may require longer solution time than metaheuristic, it allows to get the optimal solutions for several problems, and to verify the efficiency of the metaheuristic methods.

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| Cell Formation Problem Solved Exactly with the Dinkelbach Algorithm |
|---------------------------------|------------------|--------|--------|-----------------|--------|
| P15 Srinivasan et al. [42]      | 16               | 30     | 6      | 69.53           | 69.53  |
| P16 King [43]                   | 16               | 43     | 8      | 57.53           | 57.53  |
| P17 Carrie [44]                 | 18               | 24     | 9      | 57.73           | 57.73  |
| P18 Mosier and Taube [45]       | 20               | 20     | 5      | 43.45           | 43.06  |
| P19 Kumar et al. [46]           | 20               | 23     | 7      | 50.81           | 50.81  |
| P20 Carrie [44]                 | 20               | 35     | 5      | 77.91           | 77.91  |
| P21 Boe and Cheng [47]          | 20               | 35     | 5      | 57.98           | 57.98  |
| P22 Chandrasekharan Rajagopalon [48] | 24           | 40     | 7      | 100             | 100    |
| P23 Chandrasekharan Rajagopalon [48] | 24           | 40     | 7      | 85.11           | 85.11  |
| P24 Chandrasekharan Rajagopalon [48] | 24           | 40     | 7      | 73.51           | 73.51  |
| P25 Chandrasekharan Rajagopalon [48] | 24           | 40     | 11     | 53.29           | 53.29  |
| P26 Chandrasekharan Rajagopalon [48] | 24           | 40     | 12     | 48.95           | 48.95  |
| P27 Chandrasekharan Rajagopalon [48] | 24           | 40     | 12     | 47.26           | 47.26  |
| P28 McCormick [41]              | 27               | 27     | 5      | 54.82           | 54.82  |
| P29 Carrie [44]                 | 28               | 46     | 10     | 47.06           | 47.06  |
| P30 Kumar and Vannelli [49]     | 30               | 41     | 14     | 63.31           | 63.31  |
| P31 Stanfel [40]                | 30               | 50     | 13     | 60.12           | 60.12  |
| P32 Stanfel [40]                | 30               | 50     | 14     | 50.83           | 50.83  |
| P33 King and Nakornchai [28]    | 36               | 90     | 17     | 47.75           | 47.75  |
| P34 McCormick [41]              | 37               | 53     | 3      | 60.64           | 60.64  |
| P35 Chandrasekharan Rajagopalon [13] | 40           | 100    | 10     | 84.03           | 84.03  |

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5 Conclusion

The cell formation problem can be transformed into an equivalent binary linear fractional programming. We introduce a solution approach based on the Dinkelbach algorithm where the CPLEX 12.2 Optimizer is used to solve the binary linear programming problem at each iteration. The optimal value of the grouping efficacy $Eff$ is obtained for 27 out of the 35 benchmarked problems often used in the literature. The best known solution of each of these 27 problems is equal to the optimal value of $Eff$. Hence our exact procedure may require longer solution time, but it allows to solve exactly several benchmarked problems, and to show the efficiency of the metaheuristic reaching the best known solutions.

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References


