Enumeration of Benzenoid and Coronoid Hydrocarbons

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This is a report on the present status of computer-aided enumeration and classification of benzenoids and coronoids (polyhexes). The existing data from previous works are displayed in comprehensive tables. Numerous data from papers in press are included, as well as original contributions.

Introduction

The problem of "cell-growth" is a classical problem in mathematics [1-5]. When applied to hexagonal "animals" it has relevance to the studies of benzenoid (polycyclic aromatic) hydrocarbons, especially to their enumeration [6-27]. The research field is rapidly expanding, as is documented by the numerous papers in press or just appeared [28-34].

In works of this type it is important to define precisely the classes of systems which are subjected to enumeration. Here a benzenoid is defined in consistence with a review [35] and a book [36]. It is a planar system of identical regular hexagons, which are simply connected. The last restriction is released in the definition of coronoids. In those systems (also called "true" circulenes) a hole is recognized, which should have a size of at least two hexagons. Multiple coronoids with more than one hole are possible. It is emphasized that we adher to the definitions which exclude helicene derivatives and similar systems.

A catacondensed system (benzenoid [35] or coronoid [28]) does not possess any vertex belonging to three hexagons [3, 35-40]. The feature is manifested by the absence of triangles in the dualist

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graph. All other systems are referred to as pericondensed. A system is non-Kekuléan when it does not possess any Kekulé structure; in the opposite case it is called Kekuléan. All catacondensed systems are known to be Kekuléan. Alternative designations for catacondensed benzenoids and pericondensed benzenoids are catafusenes and perifusenes, respectively.

The term polyhex seems to be a suitable designation for benzenoids and coronoids together.

In general it is noted that no terminology in this area has been established as a prevailing standard at present.

In the present work we summarize some previous results on enumeration of benzenoids and coronoids. Also new results are reported, where the ranges of computation have been extended with regard to the h value. Here h denotes the number of hexagons of a system.

The footnotes to tables (see below) are supposed to give the credit of priority to the different researchers. The quoted numbers are either found explicitly in the cited references or obtained simply by additions or subtraction of published numbers. Still the quotations do not give full justice to all researchers which have developed this field because of the special definitions of benzenoids and coronoids adherred to here. These conventions discriminate in particular the important works of

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Harary and Read [3] and of Lunnon [4]. These authors have included helicenes not covered by the definitions of the present work. Similarly the works of Balaban and Harary [6] and Balaban [7, 8] are discriminated due to the inclusion of helicenes.

In the present report the number of hexagons, h, is the leading parameter. As a consequence it makes the extensive works of Dias [13, 14, 17–19, 22] fall outside the present scope. This researcher concentrated upon the chemical isomerism, which is reflected by the chemical formula C_nH_s . For polyhexes n is the total number of vertices and s the number of vertices of second degree. The works of Cyvin [24] and Cyvin et al. [26, 31, 32], where the benzenoids are enumerated according to their Kekulé structure counts, fall also outside the scope of the present report.

Benzenoids

Catacondensed Benzenoids

Table 1 shows the results of enumeration for catacondensed benzenoids (catafusenes). The unbranched systems are classified according to their symmetries into [6]:

```
a acenes (linear): D_{6h} for h = 1, otherwise D_{2h}; 
m mirror-symmetrical, C_{2r}; 
c centrosymmetrical, C_{2h}; 
u unsymmetrical, C_s.
```

Pericondensed Benzenoids

Table 2 shows the results of enumeration for pericondensed benzenoids (perifusenes). All non-Kekuléans (*o*) are pericondensed. The Kekuléan pericondensed systems are divided into:

np normal pericondensed;e essentially disconnected [41].

The latter systems (e) possess edges which have fixed bonds (simple or double) in all Kekulé structures. All other Kekuléans are referred to as normal [23, 25].

Coronoids (Single)

Multiple coronoids exist, the smallest one having h = 13 [14]. They are not counted in this report.

Table 1. Numbers of	of catacondensed	benzenoids	(catafusenes).
---------------------	------------------	------------	----------------

h	Unbr	anched			Total unbranched	Branched	Total catacondensed
	а	m	С	и			
1	1 a	0	0	0	1 ^a	0	1 ^a
2	1 a	0	0	0	1 ^a	0	1 a
3	1 a	1 ^a	0	0	2 a	0	2 a
4	1 a	1 a	1 a	1 a	4 a	1 a	5 a
5	1 a	4 a	1 a	4 a	10 ^a	2 a	12 a
6	1 a	3 b	4 a	16 a	24 °	12°	36 ^d
7	1 a	12 ^b	4 a	50 b	67°	51 °	118 ^d
8	1 a	10 ^b	13 a	158 b	182 °	229°	411 ^d
9	1 b	34 b	13 ^b	472 b	520 b, e	969 b, e	1 489 ^d
10	1 b	28 b	39 b	1 406	1 474	4 098	5 572 d
11	1 b	97 ^b	39 ^b	4 111	4 248	16 867	21 115 f
12	1 b	81 b	116 ^b	11 998	12 196	†	†
13	1	271	115	34 781	35 168	†	†
14	1	226	339	100 660	101 226	†	†
15	1	764	336	290 464	291 565	†	†
16	1	638	988	837 137	838 764	†	†
17	1	2 141	977	2 408 914	2 412 033	†	†
18	1	1 787	2 866	6 925 100	6 929 754	†	†
19	1	6 025	2 832	19 888 057	19 896 915	†	†
20	1	5 030	8 298	57 071 610	57 084 939	†	+

^a Balaban and Harary [6]. – ^b Brunvoll, Cyvin, and Cyvin [29]. – ^c He and He [20]. – ^d Knop, Szymanski, Jeričević, and Trinajstić [11]. – ^e He and He [33]. – ^f Stojmenović, Tošić, and Doroslovački [34]. – [†] Unknown.

Table 2. Numbers of pericondensed benzenoids (perifusenes).

h	Kekuléan		Total pericon-	Non- Kekuléan	Total pericon-	
	np	e	densed Kekuléan*	Rekulcan	densed **	
3	0	0	0	1 a	1	
4	1 a	0	1	1 a	2	
4 5	2 b	1 b	3	7 a	10	
	12 b	3 b	15	30 a	45	
6 7	49 b	23 b	72	141 c	213	
8	232 b	121 b	353	671 c	1 024	
9	1 042 b	692 b	1 734	3 282 b	5 016	
10	4 803 ^d	3 732	8 535	15 979	24 514	
11	21 804 d	19 960	41 764	78 350	120 114	

* The sums of numbers in the two first columns.

** Obtained as differences from published data (Grand

total – Total catacondensed).

^a Balaban and Harary [6]. – ^b Brunvoll, Cyvin, and Cyvin [29]. – ^c He and He [20]. ^d Cyvin, Brunvoll, and Cyvin [32].

Table 3. Numbers of catacondensed coronoids.

h	Primitive (unbranched)	Annelated (branched)	Total catacondensed *
8	1 a	0	1
9	1 a	2 a	3
10	3 a	12 a	15
11	2 a	60 a	62
12	11 a	301	312
13	12 a	1423	1435
14	40 a	†	†
15	68	Ť	†
16	192	†	†
17	395	†	†
18	1061	†	Ť
19	2432	†	†
20	6318	†	†

The sums of numbers of the two first columns.

Brunvoll, Cyvin, and Cyvin [28]. - † Unknown.

Table 4. Numbers of pericondensed coronoids.

h	Kekulé	ean	Total peri- condensed		Total		
	np	e	Kekuléan*	Rekuleali	pericon- densed **		
9	0	0	0	2 a	2 a		
10	9 a	0	9	19 a	28 a		
11	64 a	2 a	66	155 a	221 a		
12	508	34	542	1100 a	1642		

The sums of numbers in the two first columns.

Sum Pericondensed Kekuléan + Non-Kekuléan.

Brunvoll, Cyvin, and Cyvin [28].

Table 3 shows the numbers of catacondensed coronoids. Primitive [28] systems are defined as consisting of one single circular chain. The other catacondensed systems have at least one branching hexagon and are sometimes referred to as annelated.

Table 4 shows the number of pericondensed coronoids. The designations np and e refer to normal pericondensed and essentially disconnected systems in analogy with the benzenoids (see above, especially Table 2).

The "neo" Classification

The usefulness of a distinction between normal (n) and essentially disconnected (e) benzenoids or coronoids among the Kekuléan systems has been pointed out previously [24, 26, 29]. Here the normal benzenoids or coronoids refer to the catacondensed systems together with the normal pericondensed. The inclusion of non-Kekuléan systems (o) gives the grand total (n + e + o). The concept neo classification refers to the n, e and o classes. The different numbers of the benzenoids and coronoids (in parentheses) within this classification are summarized in Table 5.

More Special Classifications

Symmetry

The classification of unbranched catacondensed benzenoids according to symmetry was first undertaken by Balaban and Harary [6], and pursued by Brunvoll et al. [29]; cf. Table 1. It has been pointed out that the benzenoids in general can belong to one of the eight symmetries [42]: D_{6h} , C_{6h} , D_{3h} , C_{3h} , D_{2h} , C_{2h} , C_{2v} or C_s . The same is the case for coronoids. Table 6 summarizes the distribution into the different symmetries for all benzenoids (also pericondensed and branched catacondensed) and coronoids; the latter results are given in parentheses.

△ Values

The positive integer Δ is an important parameter for benzenoids and coronoids. It is defined as the absolute magnitude of the difference between the numbers of black and white vertices, referring to

Table 5. The "neo" classification for benzenoids and coronoids (numbers in parentheses).

h	n	e	o*	Total benzenoids (coronoids)	Total benzenoids + coronoids
1	1 a	0	0	1 b	1 °
2	1 a	0	0	1 b	1 °
3	2 a	0	1 ^d	3 b	3 c
4	6 a	0	1 ^d	7 b	7°
5	14 ^a	1 a	7 ^d	22 b	22°
6	48 a	3 a	30 ^d	81 e	81 °
7	167 a	23 a	141 ^f	331 e	331 °
8	643 ^a (1 ^g)	121 a	671 ^f	1 435 e (1 h)	1 436 °
9	2 531 a (3 g)	692 a	$3282^{a}(2^{g})$	6 505 e (5 h)	6 510 c, i
10	$10\ 375^{\mathrm{j}}(24^{\mathrm{g}})$	3 732	15979(19g)	30 086 e (43 h, k)	30 129 i
11	42 919 ^j (126 ^g)	$19960(2^{g})$	78 350 (155 g)	141 229 ¹ (283 g)	141 512
12	† (820)	† (34)	$†(1100^{\circ})$	† (1954 ^g)	†

Also Table 2 (coronoids Table 4).

Balaban and Harary [6].

He and He [20].

Cioslowski [30].

Cyvin, Brunvoll, and Cyvin [32].

Table 6. Numbers of benzenoids and coronoids (in parentheses) of different symmetries.

h	D_{6h}	C_{6h}	D_{3h}	C_{3h}	D_{2h}	C_{2h}	C_{2v}	C_s
1	1	0	0	0	0	0	0	0
2	0	0	0	0	1	0	0	0
3	0	0	1	0	1	0	1	0
4	0	0	1	0	2	1	1	2
5	0	0	0	0	2	1	9	10
6	0	0	1	1	3	7	12	57
7	1	0	1	1	3	7	39	279
8	0	0	0	0	6(1)	35	61	1 333
9	0	0	1(1)	5	7 (0)	36	178 (1)	6 278 (3)
0	0	0	4(0)	5	11 (3)	169 (3)	274 (9)	29 623 (28)
1	0	0	0 (0)	0	14(0)	177 (0)	796 (24)	140 242 (259)
2	0(1)	0	3 (3)	21 (2)	21 (10)	807 (25)	† (79)	† (1834)
3	2(0)	0	4(0)	26 (0)	23 (0)	859 (0)	† (185)	† (†)
4	0(0)	0	0(0)	0(0)	41 (†)	† (†)	† (†)	† (†)
5	0(0)	0	3 (9)	95 (17)	50 (0)	† (0)	† (†)	† (†)
6	0(0)	0	12 (0)	118 (0)	80 (†)	† (†)	† (†)	† (†)
7	0(0)	0	0 (0)	0(0)	94 (0)	† (0)	† (†)	† (†)
8	0 (3)	0(1)	6 (23)	423 (102)	156 (†)	† (†)	† (†)	† (†)
19	2 (0)	2(0)	19 (0)	543 (0)	189 (0)	† (0)	† (†)	† (†)
20	0(0)	0(0)	0 (0)	0 (0)	310 (†)	† (†)	† (†)	† (†)

[†] Unknown.

the coloring of vertices ("black and white" is synonymous with "starred and unstarred") [37, 40]. This number (Δ) is also equal to the absolute magnitude of the difference between the numbers of valleys and peaks [43, 44]. Furthermore, Δ is also equal to the absolute magnitude of the difference between the numbers of upright and upset triangles [20, 45] in the dualist graph [6]. It is long known that all Kekuléan systems have $\Delta = 0$, and that this is not a sufficient condition [37]. Non-Kekuléan benzenoids and coronoids with $\Delta = 0$ exist; benzenoids when $h \ge 11$ [37] and coronoids when $h \ge 15$ [28]. They are called concealed non-Kekuléans [28, 44]. The others $(\Delta > 0)$ are referred to as obvious non-Kekuléan [44]. Table 7 shows the distribution of △ values for all benzenoids (and coronoids).

Brunvoll, Cyvin, and Cyvin [29]. Klarner [1].

He and He [33].

Knop, Szymanski, Jeričević, and Trinajstić [11].

Brunvoll, Cyvin, and Cyvin [28].

Knop, Szymanski, Jeričević, and Trinajstić [16].

Correction: Knop, Müller, Szymanski, and Trinajstić

^{[25].} Stojmenović, Tošić, and Doroslovački [34]. Unknown.

Table 7. Numbers of benzenoids and coronoids (in parentheses) with different Δ values.

h	$\Delta = 0*$	$\Delta = 1$	$\Delta = 2$	$\Delta = 3$	$\Delta = 4$
1	1	0	0	0	0
2	1	0	0	0	0
3	. 2	1 a	0	0	0
4	6	1 a	0	0	0
5	15	7 a	0	0	0
6	51	28 a	2 a	0	0
7	190	134 a	7 a	0	0
8	$764(1^{b})$	619 a	52 a	0	0
9	$3223(3^{b})$	$2957(2^{b})$	322	3	0
10	$14107(24^{b})$	14024(16 ^b)	$1916(3^{b})$	39	0
11	62887(128 ^b)	67 046(131 ^b)	10922(23 ^b)	$374(1^{b})$	0
12	†(854 b)	$†(906^{b})$	†(182 ^b)	$†(12^{b})$	14

^{*} For benzenoids when $h \le 10$: All Kekuléan, viz. catacondensed (Table 1) + np + e (Table 2). All coronoids when $h \le 14$ are Kekuléan, viz. catacondensed (Table 3) + np + e (Table 4).

+ np + e (Table 4). ^a Brunvoll, Cyvin, and Cyvin [29]. – ^b Brunvoll, Cyvin, and Cyvin [28]. – [†] Unknown.

It was demonstrated by an inductive reasoning that the maximal Δ value for a benzenoid is

$$\Delta_{\max} = \left[\frac{h}{3} \right],$$

where [x] is used to designate the largest integer smaller than or equal to x.

Numbers of Internal and External Vertices

The computer program of Knop and Trinajstić with collaborators [11, 12, 16, 21] is based on the fact that a benzenoid is uniquely defined by its perimeter. For the purpose of identification these authors devised a boundary code given in terms of a six-letter alphabet, actually consisting of a sequence of integers. Basically the same idea was first used by Balaban [46], and was employed with modifications by Doroslovački and Tošić with collaborators [15, 23, 34].

Table 8 includes the data from Knop et al. [11] concerning the classification of benzenoids according to their number of internal vertices, say n_i .

Stojmenović et al. [34] performed a classification according to the length of the perimeter. Actually their material contains the enumeration of all benzenoids with the perimeter length up to 46 edges and classified according to the h values. The length of the perimeter amounts to the same as the number of external vertices, say $n_{\rm e}$. This number, being one of the invariants of a benzenoid, is closely related to

the invariant n_i . From the known connections between invariants of a benzenoid [35, 38] it is obtained

$$n_i = 2h - (n_e/2) + 1$$
.

Hence also the data of Stojmenović et al. [34] contain the information about the numbers of internal vertices. As such most of these data are entered into Table 8. The table would be too voluminous if all the available data [34] were included. They cover all benzenoids up to $n_e = 46$, which makes the highest entry of h equal to 44. Actually a unique benzenoid with h = 44 and $n_e = 46$ is listed; it has 66 internal vertices ($n_i = 66$).

The open spaces to the right and above the figures in Table 8 are equivalent to zeros. They reflect the fact that n_i has a maximal value for a given h. According to Harary and Harborth [5] this maximum is

$$(n_i)_{\text{max}} = 2h + 1 - \{(12h - 3)^{1/2}\},\,$$

where $\{x\}$ is used to denote the smallest integer greater than or equal to x. We also deduce straightforwardly that the length of the perimeter is bounded as

$$2\{(12h-3)^{1/2}\} \le n_e \le 4h+2.$$

As an example, the formulas give for h = 44: $(n_i)_{\text{max}} = 66$, $(n_e)_{\text{min}} = 46$, both figures in consistence with the above quoted case.

The ABCDEF Classification

He and He [20, 33] have classified the benzenoids and coronoids into the following six classes. A are the Kekuléans, B+C cover the catacondensed benzenoid systems: B unbranched and C branched. D+F cover the pericondensed benzenoids, D not possessing any end hexagon and F possessing at least one end hexagon, i.e. one with five free edges and manifested by a vertex of degree one in the dualist graph. The class E is coincident with the primitive coronoids. The rest of the coronoids come under D and F depending on whether they have not or have end hexagons, respectively. The classes D and F are separated into Kekuléans and non-Kekuléans. It is referred to Table 9 for enumeration results.

This classification is conveniently performed by means of (a) Δ values equal to |U-D| and (b) (U+D) values, where U and D are the respective numbers of upright and upset triangles in the

Table 8. Numbers of benzenoids with different numbers of internal vertices, n_i .

h	$n_i = 0*$	1		2		3	4	5		6	7		8	9	10	11	12	13
1	1																	
2 3 4 5 6	2	1 a																
4	2 5	1 a		1 a														
5	12	6 a		3 a		1 a	2.0											
7	36 118	24 ^a 106 ^a		14 ^a 68 ^a		4 a 5 a	3 a 10 a	3	a	1 a								
8	411	453 a		329 a	144		67 a	21		9 a	1:	a						
9	1 489	1 966 a		601 a	82.	5 a	396 a	154		55 a	15		4 a					
10	5 572	8 395 a		652 a	4 518		2 340 a	1 018		416a	123		42 a	9 a		a a b		
11	21 115	35 885 b		109 ^b 318 ^b	24 020 124 119		13 415 ^b 74 985 ^b	6 405 38 727		811 b 306 b	1 008 7 375		333 ^b 713 ^b	100 b 878 b			13 ^b	1 b
12		1	108	3100	124 113	9 °	14 983 0	30 121	10	300 °	1313		/15	0/0	219	01	13-	1 -
h	$n_i = 4$		5	6		7	8	9	9	10	11		12		13	14		15
13	408 785	226 8		4 326			20 119	7 25		2 459	708		187		35	4		
14	1		† 69	1 933	328 5		141 268	55 124		0 321	6 594		2 001		33	120 1 570		16
15 16	1		Ţ	Ţ		Ţ	947 291	395 86		5 656	55 857 436 698		8 396 6 434	5 6 51 6		16 234		347 501
17	+	-	+	+		+	+		† 115.	†	†30 076		6 839			148 430		166
18	i	-	†	†		†	†		t	†	†		†		† 1	262 442	424	429
h	$n_{\rm i} = 16$	17		18	19		20	21	22		23	24	2	25	26	27	r.	28
14	1																	
15	70	4		20														
16 17	1 121 13 286	223 3 414		30 763	117		9											
18	133 713	39 143	10	587	2 437		471	53	3									
19		388 180	116	648	32 042		7 885	1 647	256		18	1						
20	†	†	1 140	529 3	345 834			25 050	5 726			129		4	4.7			
21	Ţ	Ţ		Ţ	Ţ	1 02	29 521 2		79 472 03 415			838 027	58 13 65		47 2 467	290		16
22 23	+	+		Ť	+		Ţ	† 9	03 415	247	989 62 † 771		13 03 200 54		7 167	9 349		16 448
24	+	+		+	+		+	+	+		† //1	†	200 5-			157 989		324
25	÷	†		†	†		Ť	†	+		†	†		†	†	†		218

Table 9. Numbers of benzenoids and coronoids (numbers in parentheses) according to the ABCDEF classification.

h	A Kekuléan					Non-Kekuléar	1
	B*	C*	D	E**	F	D	F
1	1	0	0	0	0	0	0
2	1	0	0	0	0	0	0
3	2	0	0	0	0	1 a	0
4	4	1	1 a	0	0	0 a	1 a
5	10	2	1 a	0	2 a	1 a	6 a
6	24	12	3 a	0	12 a	3 a	27 a
7	67	51	6 a	0	66 a	9 a	132 a
8	182	229	22 a	(1)	331 a	27 a	644 a
9	520	969	60 b	(1)	1 674 (2)	117(2)	3 165 b
0	1 474	4 098	224 (9)	(3)	8 311 (12)	437 (6)	15 542 (13)
1	4 248	16 867	806 (13)	(2)	40 958 (113)	1 778 (52)	76 572 (103

 ^{*} Catacondensed benzenoids; see Table 1. - † Unknown.
 a Knop, Szymanski, Jeričević, and Trinajstić [11].
 b Stojmenović, Tošić, and Doroslovački [34]. All data for h > 10 are taken from this reference.

^{*} See Table 1 (catacondensed); B unbranched, C branched. ** See Table 3 (primitive). — ^a He and He [20]. — ^b He and He [33].

dualist graph, and (c) V_1 , V_2 and V_3 , designating numbers of vertices of the dualist graph with degree one, two and three, respectively.

These values are easily obtained from the Hematrix [20]. It is noted that in the graph generation program of He and He [20], considering the connectivity of the graph and using the properties of the He-matrix, the coronoids are automatically included into the classification. In particular the He-matrix can be used to classify the systems according to Δ values. A He-matrix is also suited for the studies of the rotation and reflection transformations. Thus it may be used to determine the symmetry groups of the benzenoid and coronoid systems.

Estimates for the Number of Benzenoids

As the number h of hexagons increases, the direct enumeration of benzenoids becomes extremely difficult and even with the use of the most powerful computing devices one cannot expect the solution of this task for the totality of benzenoids with, say, h = 20 or h = 30. Another approach to this problem would be the application of Pólya's theory, which provides general mathematical techniques for enumerations of combinatorial objects of various types [47]. Unfortunately, the enumeration of benzenoids by means of Pólya's method could not be realized until now and is considered as an open problem of combinatorics. From a mathematician's point of view, the enumeration of benzenoid systems is an unsolved problem.

If helicenes and other non-planar systems are not disregarded, then for large values of h an exact asymptotic behaviour of the number of catacondensed benzenoids is known [3] and has the form

$$N \approx a h^p b^h$$
.

where p, a and b are specified constants. This result does not apply for catacondensed systems as defined in the present paper. Nevertheless, it has been attempted [27] to reproduce an approximate asymptotic behaviour of catacondensed benzenoids with helicenetype systems disregarded by the same mathematical form, but other values of the parameters p, a and b. By a fitting procedure, using the known numbers for $h \le 11$, Gutman [27] determined p = -5/4, a = 0.049 and b = 4.27. In an analogous way he also determined a set of parameters, viz. p = -3/2, a = 0.045

Table 10. Comparison between exact and estimated numbers of catacondensed benzenoids and all benzenoids.

h	Catacono	densed benzenoids	Total ben	zenoids
	Exact	Estimate ^a	Exact	Estimate b
6	36	32	81	76
7	118	111	331	$\frac{325}{\pm 10}$
8	411	402	1 435	1 440 ± 40
9	1 489	1 480 ± 30	6 505	6 510 ± 180
10	5 572	5 550 ± 120	30 086	30 000 ± 900
11	21 115	21 050 ± 500	141 229	140 400 ± 4 400
12	†	$ \begin{array}{r} $	†	666 000 ± 22 000
13	†	$\begin{array}{c} \pm 2000 \\ 311000 \\ \pm 8000 \end{array}$	†	3 190 000 ± 110 000
14	†	$\begin{array}{c} \pm \ 0000 \\ 1\ 212\ 0000 \\ \pm \ 32\ 0000 \end{array}$	†	15 400 000 ± 570 000
15	†	$\begin{array}{c} \pm 32000 \\ 4750000 \\ \pm 130000 \end{array}$	†	$75\ 000\ 000$ $\pm 2\ 900\ 000$

^a The constants a and b (see the text for definitions) are assumed to have the values 0.049 ± 0.0005 and 4.27 ± 0.005 , respectively. – ^b $a = 0.045 \pm 0.0005$, $b = 5.4 \pm 0.01$.

† Unknown.

and b = 5.4, for the total number of benzenoids (both catacondensed and pericondensed). A corresponding analysis for normal benzenoids is due to Cyvin et al. [32].

In Table 10 the exact and estimated numbers of catacondensed benzenoids and benzenoids in total are given for $6 \le h \le 15$.

Conclusion

The present report does not imply an end to the enumeration and classification of polyhexes. On the contrary, the activity in this research area has apparently been intensivated during the last years, and not least during the preparation of this report. New results are to be awaited, further extensions to higher h values on one hand and new types of classifications on the other.

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- [1] D. A. Klarner, Fibonacci Quarterly 3, 9 (1965).
- [2] F. Harary, in: Graph Theory and Theoretical Physics (F. Harary, ed.). Academic Press, London 1967; p. 1.
- [3] F. Harary and R. C. Read, Proc. Edinburgh Math. Soc. 17 (Ser. II), 1 (1970).
- [4] W. F. Lunnon, in: Graph Theory and Computing (R. C. Read, ed.). Academic Press, New York 1972, p. 87.
- [5] F. Harary and H. Harborth, J. Combinat. Inf. & System Sci. 1, 1 (1976).
- [6] A. T. Balaban and F. Harary, Tetrahedron 24, 2505 (1968).
- [7] A. T. Balaban, Tetrahedron 25, 2949 (1969).
- [8] A. T. Balaban, Match (Mülheim) 2, 51 (1976)
- [9] K. Balasubramanian, J. J. Kaufman, W. S. Koski, and A. T. Balaban, J. Comput. Chem. 1, 149 (1980).
- [10] J. R. Dias, J. Chem. Inf. Comput. Sci. 22, 15 (1982).
- [11] J. V. Knop, K. Szymanski, Ž. Jeričević, and N. Trinajstić, J. Comput. Chem. **4**, 23 (1983).
- [12] N. Trinajstić, Ž. Jeričević, J. V. Knop, W. R. Müller, and K. Szymanski, Pure & Appl. Chem. 55, 379 (1983).
- [13] J. R. Dias, Can. J. Chem. 62, 2914 (1984).
- [14] J. R. Dias, J. Chem. Inf. Comput. Sci. 24, 124 (1984).
- [15] R. Doroslovački and R. Tošić, Rev. Res. Fac. Sci. Univ. Novi Sad, Math. Ser. 14, 201 (1984).
- [16] J. V. Knop, K. Szymanski, Ž. Jeričević, and N. Trinajstić, Match (Mülheim) 16, 119 (1984).
- [17] J. R. Dias, Accounts Chem. Res. 18, 241 (1985).
- [18] J. R. Dias, J. Macromol. Sci. Chem. A 22, 335 (1985).
- [19] J. R. Dias, Nuov. J. Chim. 9, 125 (1985)
- [20] W. He and W. He, Theor. Chim. Acta **68**, 301 (1985).
- [21] J. V. Knop, W. R. Müller, K. Szymanski, and N. Trinajstić, Computer Generation of Certain Classes of Molecules, Union of Chemists and Technologists of Croatia, Zagreb 1985.
- [22] J. R. Dias, J. Mol. Struct. (Theochem.) 137, 9 (1986).
- [23] R. Tošić, R. Doroslovački, and I. Gutman, Match (Mülheim) 19, 219 (1986).
- [24] S. J. Cyvin, Match (Mülheim) 20, 165 (1986).
- [25] J. V. Knop, W. R. Müller, K. Szymanski, and N. Trinajstić, Match (Mülheim) 20, 197 (1986).

- [26] S. J. Cyvin and I. Gutman, Z. Naturforsch. **41a**, 1079 (1986).
- [27] I. Gutman, Z. Naturforsch. 41 a, 1089 (1986)
- [28] J. Brunvoll, B. N. Cyvin, and S. J. Cyvin, J. Chem. Inf. Comput. Sci. 27, 14 (1987).
- [29] J. Brunvoll, S. J. Cyvin, and B. N. Cyvin, J. Comput. Chem. 8, 189 (1987).
- [30] J. Cioslowski, J. Comput. Chem. (in press).
- [31] B. N. Cyvin, J. Brunvoll, S. J. Cyvin, and I. Gutman, Match (Mülheim) 21, 301 (1986).
- [32] S. J. Cyvin, J. Brunvoll, and B. N. Cyvin, Z. Naturforsch. 41a, 1429 (1986).
- [33] W. He and W. He, Tetrahedron **42**, 5291 (1986).
- [34] I. Stojmenović, R. Tošić, and R. Doroslovački, in: Graph Theory, Proceedings of the Sixth Yugoslav Seminar on Graph Theory Dubrovnik, April 18–19, 1985 (R. Tošić, D. Acketa, and V. Petrović, eds). University of Novi Sad, Novi Sad 1986, p. 189.
- [35] I. Gutman, Bull. Soc. Chim. Beograd 47, 453 (1982).
- [36] I. Gutman and O. E. Polanski, Mathematical Concepts in Organic Chemistry. Springer-Verlag, Berlin 1986.
- [37] I. Gutman, Croat. Chem. Acta 46, 209 (1974).
- [38] O. E. Polansky and D. H. Rouvray, Match (Mülheim) **2,** 63 (1976).
- [39] O. E. Polansky and D. H. Rouvray, Match (Mülheim) 3, 97 (1977).
- [40] N. Trinajstić, Chemical Graph Theory, Vols. I, II, CRC Press, Boca Raton, Florida 1983.
- [41] S. J. Cyvin, B. N. Cyvin, and I. Gutman, Z. Naturforsch. 40 a, 1253 (1985).
- [42] S. J. Cyvin and I. Gutman, Comp. Maths. Appls. 12 B, 859 (1986).
- [43] I. Gutman and S. J. Cyvin, J. Mol. Struct. (Theochem.) 138, 325 (1986).
- [44] S. J. Cyvin and I. Gutman, J. Mol. Struct. (Theochem.) 150, 157 (1987).
- [45] A. T. Balaban, Rev. Roum. Chim. 26, 407 (1981).
- [46] A. T. Balaban, Tetrahedron 27, 6115 (1971).
- [47] F. Harary and E. P. Palmer, Graphical Enumeration, Academic Press, New York 1973.