

ChemmineR: Cheminformatics Toolkit for R

Yiqun Cao, Tyler Backman, Kevin Horan, Thomas Girke

Email contact: thomas.girke@ucr.edu

March 24, 2014

1 Introduction

ChemmineR is a cheminformatics package for analyzing drug-like small molecule data in R. Its latest version contains functions for efficient processing of large numbers of small molecules, physicochemical/structural property predictions, structural similarity searching, classification and clustering of compound libraries with a wide spectrum of algorithms.

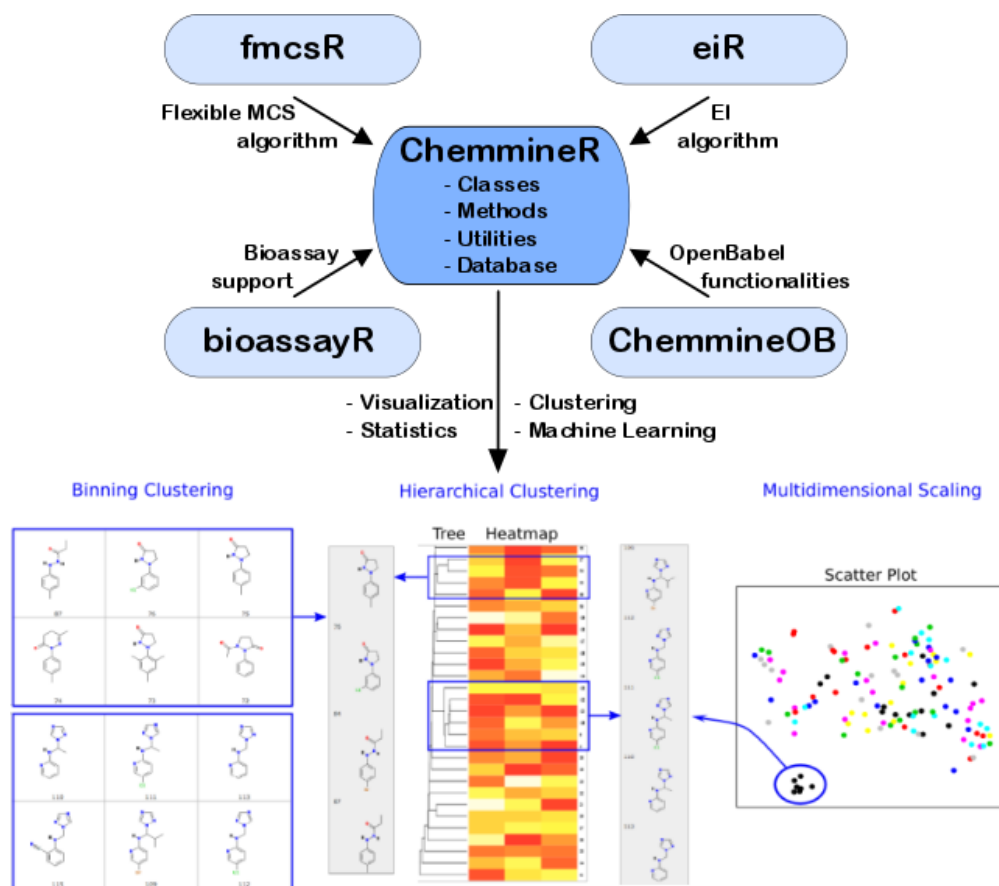


Figure 1: *ChemmineR* environment with its add-on packages and selected functionalities

In addition, *ChemmineR* offers visualization functions for compound clustering results and chemical structures. The integration of cheminformatic tools with the R programming environment has many advantages, such as easy access to a wide spectrum of statistical methods, machine learning algorithms and graphic utilities. The first version of this package was published in Cao et al. (2008). Since then many additional utilities and add-on packages have been added to the environment and many more are under development for future releases (Figure 2; (Backman et al., 2011; Wang et al., 2013)).

Contents

1	Introduction	1
2	Recently Added Features	3
3	Getting Started	3
3.1	Installation	3
3.2	Loading the Package and Documentation	3
3.3	Five Minute Tutorial	3
4	OpenBabel Functions	9
5	Overview of Classes and Functions	9
5.1	Molecular Structure Data	9
5.2	Structure Descriptor Data	10
6	Import of Compounds	10
6.1	SDF Import	10
6.2	SMILES Import	11
7	Export of Compounds	12
7.1	SDF Export	12
7.2	SMILES Export	12
8	Format Interconversions	12
9	Splitting SD Files	13
10	Streaming Through Large SD Files	13
11	Storing Compounds in an SQL Database	14
11.1	Loading Data	15
11.2	Searching	15
12	Working with SDF/SDFset Classes	16
13	Molecular Property Functions (Physicochemical Descriptors)	17
14	Bond Matrices	20
15	Charges and Missing Hydrogens	20
16	Ring Perception and Aromaticity Assignment	20
17	Rendering Chemical Structure Images	23
17.1	R Graphics Device	23
17.2	Online with ChemMine Tools	26
18	Similarity Comparisons and Searching	28
18.1	Maximum Common Substructure (MCS) Searching	28
18.2	AP/APset Classes for Storing Atom Pair Descriptors	28
18.3	Large SDF and Atom Pair Databases	29
18.4	Pairwise Compound Comparisons with Atom Pairs	30
18.5	Similarity Searching with Atom Pairs	30
18.6	FP/FPset Classes for Storing Fingerprints	31
18.7	Atom Pair Fingerprints	32

18.8 Pairwise Compound Comparisons with PubChem Fingerprints	33
18.9 Similarity Searching with PubChem Fingerprints	33
18.10 Visualize Similarity Search Results	34
19 Clustering	34
19.1 Clustering Identical or Very Similar Compounds	35
19.2 Binning Clustering	36
19.3 Jarvis-Patrick Clustering	38
19.4 Multi-Dimensional Scaling (MDS)	41
19.5 Clustering with Other Algorithms	42
20 Searching PubChem	44
20.1 Get Compounds from PubChem by Id	44
20.2 Search a SMILES Query in PubChem	44
20.3 Search an SDF Query in PubChem	45
21 Version Information	45
22 Funding	45
23 References	45

2 Recently Added Features

- Improved SMILES support via new *SMIset* object class and SMILES import/export functions
- Integration of a subset of OpenBabel functionalities via new *ChemmineOB* add-on package (O'Boyle et al., 2011)
- Streaming functionality for processing millions of molecules on a laptop
- Mismatch tolerant maximum common substructure (MCS) search algorithm
- Fast and memory efficient fingerprint search support using atom pair or PubChem fingerprints

3 Getting Started

3.1 Installation

The R software for running ChemmineR can be downloaded from CRAN (<http://cran.at.r-project.org/>). The ChemmineR package can be installed from R using the `biocLite` install command.

```
> source("http://bioconductor.org/biocLite.R") # Sources the biocLite.R installation script.
> biocLite("ChemmineR") # Installs the package.
```

3.2 Loading the Package and Documentation

```
> library("ChemmineR") # Loads the package

> library(help="ChemmineR") # Lists all functions and classes
> vignette("ChemmineR") # Opens this PDF manual from R
```

3.3 Five Minute Tutorial

The following code gives an overview of the most important functionalities provided by *ChemmineR*. Copy and paste of the commands into the R console will demonstrate their utilities.

Create Instances of *SDFset* class:

```
> data(sdfsampl)
> sdfset <- sdfsampl
> sdfset # Returns summary of SDFset
```

An instance of "SDFset" with 100 molecules

```
> sdfset[1:4] # Subsetting of object
```

An instance of "SDFset" with 4 molecules

```
> sdfset[[1]] # Returns summarized content of one SDF
```

An instance of "SDF"

```
<<header>>
```

Molecule_Name	Source
"650001"	" -OEChem-07071010512D"
Comment	Counts_Line
" " 61 64 0 0 0 0 0 0 0 0 0999 V2000"	

```
<<atomblock>>
```

	C1	C2	C3	C5	C6	C7	C8	C9	C10	C11	C12	C13	C14	C15	C16
O_1	7.0468	0.0839	0	0	0	0	0	0	0	0	0	0	0	0	0
O_2	12.2708	1.0492	0	0	0	0	0	0	0	0	0	0	0	0	0
...
H_60	1.8411	-1.5985	0	0	0	0	0	0	0	0	0	0	0	0	0
H_61	2.6597	-1.2843	0	0	0	0	0	0	0	0	0	0	0	0	0

```
<<bondblock>>
```

	C1	C2	C3	C4	C5	C6	C7
1	1	16	2	0	0	0	0
2	2	23	1	0	0	0	0
...
63	33	60	1	0	0	0	0
64	33	61	1	0	0	0	0

```
<<datablock>> (33 data items)
```

PUBCHEM_COMPOUND_CID	PUBCHEM_COMPOUND_CANONICALIZED	PUBCHEM_CACTVS_COMPLEXITY
"650001"	"1"	"700"
PUBCHEM_CACTVS_HBOND_ACCEPTOR		
"7"	"..."	

```
> view(sdfset[1:4]) # Returns summarized content of many SDFs, not printed here
> as(sdfset[1:4], "list") # Returns complete content of many SDFs, not printed here
```

An SDFset is created during the import of an SD file:

```
> sdfset <- read.SDFset("http://faculty.ucr.edu/~tgirke/Documents/
+ R_BioCond/Samples/sdfsampl.sdf")
```

Miscellaneous accessor methods for SDFset container:

```
> header(sdfset[1:4]) # Not printed here
```

```
> header(sdfset[[1]])
```

```

Molecule_Name
"650001"
Source
"-OEChem-07071010512D"
Comment
"" " 61 64 0 0 0 0 0 0 0 0 0999 V2000"
Counts_Line

```

```
> atomblock(sdfset[1:4]) # Not printed here
```

```
> atomblock(sdfset[[1]])[1:4,]
```

```

      C1      C2 C3 C5 C6 C7 C8 C9 C10 C11 C12 C13 C14 C15 C16
0_1  7.0468 0.0839 0 0 0 0 0 0 0 0 0 0 0 0 0
0_2 12.2708 1.0492 0 0 0 0 0 0 0 0 0 0 0 0 0
0_3 12.2708 3.1186 0 0 0 0 0 0 0 0 0 0 0 0 0
0_4  7.9128 2.5839 0 0 0 0 0 0 0 0 0 0 0 0 0

```

```
> bondblock(sdfset[1:4]) # Not printed here
```

```
> bondblock(sdfset[[1]])[1:4,]
```

```

      C1 C2 C3 C4 C5 C6 C7
1  1 16  2  0  0  0  0
2  2 23  1  0  0  0  0
3  2 27  1  0  0  0  0
4  3 25  1  0  0  0  0

```

```
> datablock(sdfset[1:4]) # Not printed here
```

```
> datablock(sdfset[[1]])[1:4]
```

```

      PUBCHEM_COMPOUND_CID PUBCHEM_COMPOUND_CANONICALIZED PUBCHEM_CACTVS_COMPLEXITY
      "650001" "1" "700"
PUBCHEM_CACTVS_HBOND_ACCEPTOR
      "7"

```

Assigning compound IDs and keeping them unique:

```
> cid(sdfset)[1:4] # Returns IDs from SDFset object
```

```
[1] "CMP1" "CMP2" "CMP3" "CMP4"
```

```
> sdfid(sdfset)[1:4] # Returns IDs from SD file header block
```

```
[1] "650001" "650002" "650003" "650004"
```

```
> unique_ids <- makeUnique(sdfid(sdfset))
```

```
[1] "No duplicates detected!"
```

```
> cid(sdfset) <- unique_ids
```

Converting the data blocks in an *SDFset* to a matrix:

```
> blockmatrix <- datablock2ma(datablocklist=datablock(sdfset))
```

```
> # Converts data block to matrix
```

```
> numchar <- splitNumChar(blockmatrix=blockmatrix)
```

```
> # Splits to numeric and character matrix
```

```
> numchar[[1]][1:2,1:2] # Slice of numeric matrix
```

```

      PUBCHEM_COMPOUND_CID PUBCHEM_COMPOUND_CANONICALIZED
650001          650001          1
650002          650002          1

> numchar[[2]][1:2,10:11] # Slice of character matrix

      PUBCHEM_MOLECULAR_FORMULA
650001 "C23H28N4O6"
650002 "C18H23N5O3"
      PUBCHEM_OPENEYE_CAN_SMILES
650001 "CC1=CC(=N01)NC(=O)CCC(=O)N(CC(=O)NC2CCCC2)C3=CC4=C(C=C3)OCCO4"
650002 "CN1C2=C(C(=O)NC1=O)N(C(=N2)NCCCC)CCCC3=CC=CC=C3"

```

Compute atom frequency matrix, molecular weight and formula:

```

> propma <- data.frame(MF=MF(sdfset), MW=MW(sdfset), atomcountMA(sdfset))
> propma[1:4, ]

```

	MF	MW	C	H	N	O	S	F	Cl
650001	C23H28N4O6	456.4916	23	28	4	6	0	0	0
650002	C18H23N5O3	357.4069	18	23	5	3	0	0	0
650003	C18H18N4O3S	370.4255	18	18	4	3	1	0	0
650004	C21H27N5O5S	461.5346	21	27	5	5	1	0	0

Assign matrix data to data block:

```

> datablock(sdfset) <- propma
> datablock(sdfset[1])

```

```

$`650001`
      MF      MW      C      H      N      O      S
"C23H28N4O6" "456.4916" "23"    "28"    "4"    "6"    "0"
      F      Cl
      "0"    "0"

```

String searching in *SDFset* ():

```

> grepSDFset("650001", sdfset, field="datablock", mode="subset")
> # Returns summary view of matches. Not printed here.
> .

```

```

> grepSDFset("650001", sdfset, field="datablock", mode="index")

```

```

1 1 1 1 1 1 1 1 1
1 2 3 4 5 6 7 8 9

```

Export *SDFset* to SD file:

```

> write.SDF(sdfset[1:4], file="sub.sdf", sig=TRUE)

```

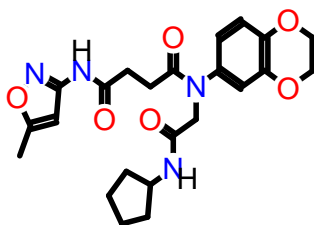
Plot molecule structure of one or many SDFs:

```

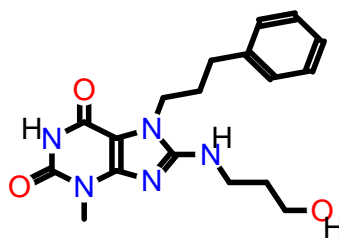
> plot(sdfset[1:4], print=FALSE) # Plots structures to R graphics device

```

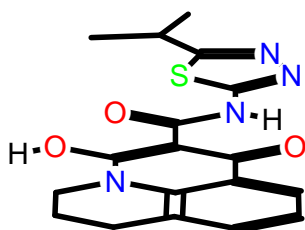
650001



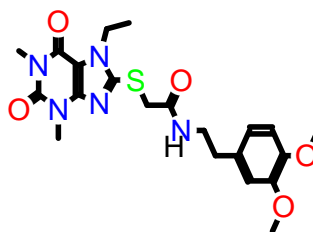
650002



650003



650004



```
> sdf.visualize(sdfset[1:4]) # Compound viewing in web browser
```

Structure similarity searching and clustering:

```
> apset <- sdf2ap(sdfset)
> # Generate atom pair descriptor database for searching
> .

> data(apset)
> # Load sample apset data provided by library.
> cmp.search(apset, apset[1], type=3, cutoff = 0.3, quiet=TRUE)

  index   cid   scores
1      1 650001 1.0000000
2     96 650102 0.3516643
3     67 650072 0.3117569
4     88 650094 0.3094629
5     15 650015 0.3010753

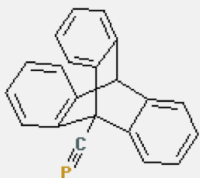
> # Search apset database with single compound.
> cmp.cluster(db=apset, cutoff = c(0.65, 0.5), quiet=TRUE)[1:4,]
```

[View Previously Accessed Compounds >>>](#)

Width of information table: 40 characters

Reference Compound (ka-01834)

[View SDF](#)[Structure Search](#)[Add to Selection](#)

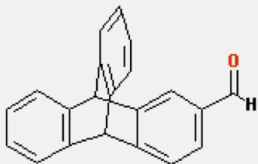


Similarities With All

ids	scores
3	0.550335570
43	0.484662577
42	0.484662577
1	0.484662577
4	0.480122324
2	0.480122324
44	0.356097561
46	0.312500000
11	0.311653117
35	0.287719298

(ChemmineR_Unnamed_Compound_3)

[View SDF](#)[Structure Search](#)[Add to Selection](#)

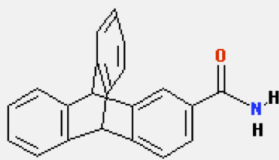


Similarities With All

ids	scores
3	1.000000000
43	0.785977860
42	0.785977860
1	0.785977860
2	0.766423358
4	0.646258503
44	0.561797753
11	0.415204678
35	0.390151515
46	0.368539326

(ChemmineR_Unnamed_Compound_43)

[View SDF](#)[Structure Search](#)[Add to Selection](#)



Similarities With All

ids	scores
43	1.000000000
42	0.840000000
1	0.840000000
3	0.785977860
2	0.709459459
4	0.611464968
44	0.601108033
11	0.390109890
46	0.339702760
35	0.223128252

Figure 2: Visualization webpage created by calling `sdf.visualize`.

8


```

sorting result...
      ids CLSZ_0.65 CLID_0.65 CLSZ_0.5 CLID_0.5
48 650049         2        48         2        48
49 650050         2        48         2        48
54 650059         2        54         2        54
55 650060         2        54         2        54

> # Binning clustering using variable similarity cutoffs.

```

4 OpenBabel Functions

ChemmineR integrates now a subset of cheminformatics functionalities implemented in the OpenBabel C++ library (O'Boyle et al., 2008, 2011). These utilities can be accessed by installing the *ChemmineOB* package and the OpenBabel software itself. *ChemmineR* will automatically detect the availability of *ChemmineOB* and make use of the additional utilities. The following lists the functions and methods that make use of OpenBabel. References are included to locate the sections in the manual where the utility and usage of these functions is described.

Structure format interconversions (see Section 8)

- `smiles2sdf`: converts from SMILES to SDF object
- `sdf2smiles`: converts from SDF to SMILES object
- `convertFormat`: converts strings between two formats
- `convertFormatFile`: converts files between two formats
- `propOB`: generates several compound properities
- `fingerprintOB`: generates fingerprints for compounds

5 Overview of Classes and Functions

The following list gives an overview of the most important S4 classes, methods and functions available in the *ChemmineR* package. The help documents of the package provide much more detailed information on each utility. The standard R help documents for these utilities can be accessed with this syntax: `?function_name` (e.g. `?cid`) and `?class_name-class` (e.g. `?SDFset-class`).

5.1 Molecular Structure Data

Classes

- *SDFstr*: intermediate string class to facilitate SD file import; not important for end user
- *SDF*: container for single molecule imported from an SD file
- *SDFset*: container for many SDF objects; most important structure container for end user
- *SMI*: container for a single SMILES string
- *SMIset*: container for many SMILES strings

Functions/Methods (mainly for SDFset container, SMIset should be coerced with smiles2sdf to SDFset)

- Accessor methods for *SDF/SDFset*
 - Object slots: `cid`, `header`, `atomblock`, `bondblock`, `datablock` (`sdfid`, `datablocktag`)
 - Summary of *SDFset*: `view`
 - Matrix conversion of data block: `datablock2ma`, `splitNumChar`
 - String search in *SDFset*: `grepSDFset`
- Coerce one class to another
 - Standard syntax `as(..., "...")` works in most cases. For details see R help with `?SDFset-class`.
- Utilities
 - Atom frequencies: `atomcountMA`, `atomcount`
 - Molecular weight: `MW`
 - Molecular formula: `MF`

- ...
- Compound structure depictions
 - R graphics device: `plot`, `plotStruc`
 - Online: `cmp.visualize`

5.2 Structure Descriptor Data

Classes

- *AP*: container for atom pair descriptors of a single molecule
- *APset*: container for many *AP* objects; most important structure descriptor container for end user
- *FP*: container for fingerprint of a single molecule
- *FPset*: container for fingerprints of many molecules, most important structure descriptor container for end user

Functions/Methods

- Create *AP/APset* instances
 - From *SDFset*: `sdf2ap`
 - From SD file: `cmp.parse`
 - Summary of *AP/APset*: `view`, `db.explain`
- Accessor methods for *AP/APset*
 - Object slots: `ap`, `cid`
- Coerce one class to another
 - Standard syntax `as(..., "...")` works in most cases. For details see R help with `?APset-class`.
- Structure Similarity comparisons and Searching
 - Compute pairwise similarities: `cmp.similarity`, `fpSim`
 - Search *APset* database: `cmp.search`, `fpSim`
- AP-based Structure Similarity Clustering
 - Single-linkage binning clustering: `cmp.cluster`
 - Visualize clustering result with MDS: `cluster.visualize`
 - Size distribution of clusters: `cluster.sizestat`

6 Import of Compounds

6.1 SDF Import

The following gives an overview of the most important import/export functionalities for small molecules provided by *ChemmineR*. The given example creates an instance of the *SDFset* class using as sample data set the first 100 compounds from this PubChem SD file (SDF): `Compound_00650001_00675000.sdf.gz` (`ftp://ftp.ncbi.nih.gov/pubchem/Compound/CURRENT-Full/SDF/`).

SDFs can be imported with the `read.SDFset` function:

```
> sdfset <- read.SDFset("http://faculty.ucr.edu/~tgirke/Documents/
+                      R_BioCond/Samples/sdfsamples.sdf")

> data(sdfsamples) # Loads the same SDFset provided by the library
> sdfset <- sdfsamples
> valid <- validSDF(sdfset) # Identifies invalid SDFs in SDFset objects
> sdfset <- sdfset[valid] # Removes invalid SDFs, if there are any
```

Import SD file into *SDFstr* container:

```
> sdfstr <- read.SDFstr("http://faculty.ucr.edu/~tgirke/Documents/
+                      R_BioCond/Samples/sdfsamples.sdf")
```

Create *SDFset* from *SDFstr* class:

```
> sdfstr <- as(sdfset, "SDFstr")
> sdfstr
```

An instance of "SDFstr" with 100 molecules

```
> as(sdfstr, "SDFset")
```

An instance of "SDFset" with 100 molecules

6.2 SMILES Import

The `read.SMiset` function imports one or many molecules from a SMILES file and stores them in a *SMiset* container. The input file is expected to contain one SMILES string per row with tab-separated compound identifiers at the end of each line. The compound identifiers are optional.

Create sample SMILES file and then import it:

```
> data(smisample); smiset <- smisample
> write.SMI(smiset[1:4], file="sub.smi")
> smiset <- read.SMiset("sub.smi")
```

Inspect content of *SMiset*:

```
> data(smisample) # Loads the same SMiset provided by the library
> smiset <- smisample
> smiset
```

An instance of "SMiset" with 100 molecules

```
> view(smiset[1:2])
```

```
$`650001`
```

An instance of "SMI"

```
[1] "O=C(NC1CCCC1)CN(c1cc2OCCOc2cc1)C(=O)CCC(=O)Nc1noc(c1)C"
```

```
$`650002`
```

An instance of "SMI"

```
[1] "O=c1[nH]c(=O)n(c2nc(n(CCCc3ccccc3)c12)NCCCO)C"
```

Accessor functions:

```
> cid(smiset[1:4])
```

```
[1] "650001" "650002" "650003" "650004"
```

```
> (smi <- as.character(smiset[1:2]))
```

```

650001
"O=C(NC1CCCC1)CN(c1cc2OCCOc2cc1)C(=O)CCC(=O)Nc1noc(c1)C"
```

```

650002
"O=c1[nH]c(=O)n(c2nc(n(CCCc3ccccc3)c12)NCCCO)C"
```

Create *SMiset* from named character vector:

```
> as(smi, "SMiset")
```

An instance of "SMiset" with 2 molecules

7 Export of Compounds

7.1 SDF Export

Write objects of classes *SDFset*/*SDFstr*/*SDF* to SD file:

```
> write.SDF(sdfset[1:4], file="sub.sdf")
```

Writing customized *SDFset* to file containing *ChemmineR* signature, IDs from *SDFset* and no data block:

```
> write.SDF(sdfset[1:4], file="sub.sdf", sig=TRUE, cid=TRUE, db=NULL)
```

Example for injecting a custom matrix/data frame into the data block of an *SDFset* and then writing it to an SD file:

```
> props <- data.frame(MF=MF(sdfset), MW=MW(sdfset), atomcountMA(sdfset))
> datablock(sdfset) <- props
> write.SDF(sdfset[1:4], file="sub.sdf", sig=TRUE, cid=TRUE)
```

Indirect export via *SDFstr* object:

```
> sdf2str(sdf=sdfset[[1]], sig=TRUE, cid=TRUE)
> # Uses default components
> sdf2str(sdf=sdfset[[1]], head=letters[1:4], db=NULL)
> # Uses custom components for header and data block
```

Write *SDF*, *SDFset* or *SDFstr* classes to file:

```
> write.SDF(sdfset[1:4], file="sub.sdf", sig=TRUE, cid=TRUE, db=NULL)
> write.SDF(sdfstr[1:4], file="sub.sdf")
> cat(unlist(as(sdfstr[1:4], "list")), file="sub.sdf", sep="\n")
```

7.2 SMILES Export

Write objects of class *SMIset* to SMILES file with and without compound identifiers:

```
> data(smisample); smiset <- smisample # Sample data set
> write.SMI(smiset[1:4], file="sub.smi", cid=TRUE)
> write.SMI(smiset[1:4], file="sub.smi", cid=FALSE)
```

8 Format Interconversions

The *sdf2smiles* and *smiles2sdf* functions provide format interconversion between SMILES strings (Simplified Molecular Input Line Entry Specification) and *SDFset* containers.

Convert an *SDFset* container to a SMILES *character* string:

```
> data(sdfsamples); sdfset <- sdfsamples[1]
> smiles <- sdf2smiles(sdfset)
> smiles
```

Convert a SMILES *character* string to an *SDFset* container:

```
> sdf <- smiles2sdf("CC(=O)OC1=CC=CC=C1C(=O)O\tname")
> view(sdf)
```

When the *ChemmineOB* package is installed these conversions are performed with the OpenBabel Open Source Chemistry Toolbox. Otherwise the functions will fall back to using the ChemMine Tools web service for this operation. The latter will require internet connectivity and is limited to only the first compound given. *ChemmineOB* provides access to the compound format conversion functions of OpenBabel. Currently, over 160 formats are supported by OpenBabel. The functions `convertFormat` and `convertFormatFile` can be used to convert files or strings between any two formats supported by OpenBabel. For example, to convert a SMILES string to an SDF string, one can use the `convertFormat` function.

```
> sdfStr <- convertFormat("SMI", "SDF", "CC(=O)OC1=CC=CC=C1C(=O)O\ttest_name")
```

This will return the given compound as an SDF formatted string. 2D coordinates are also computed and included in the resulting SDF string.

To convert a file with compounds encoded in one format to another format, the `convertFormatFile` function can be used instead.

```
> convertFormatFile("SMI", "SDF", "test.smiles", "test.sdf")
```

To see the whole list of file formats supported by OpenBabel, one can run from the command-line "obabel -L formats".

9 Splitting SD Files

The following `write.SDFsplit` function allows to split SD Files into any number of smaller SD Files. This can become important when working with very big SD Files. Users should note that this function can output many files, thus one should run it in a dedicated directory!

Create sample SD File with 100 molecules:

```
> write.SDF(sdfset, "test.sdf")
```

Read in sample SD File. Note: reading file into `SDFstr` is much faster than into `SDFset`:

```
> sdfstr <- read.SDFstr("test.sdf")
```

Run export on `SDFstr` object:

```
> write.SDFsplit(x=sdfstr, filetag="myfile", nmol=10)
> # 'nmol' defines the number of molecules to write to each file
```

Run export on `SDFset` object:

```
> write.SDFsplit(x=sdfset, filetag="myfile", nmol=10)
```

10 Streaming Through Large SD Files

The `sdfStream` function allows to stream through SD Files with millions of molecules without consuming much memory. During this process any set of descriptors, supported by *ChemmineR*, can be computed (e.g. atom pairs, molecular properties, etc.), as long as they can be returned in tabular format. In addition to descriptor values, the function returns a line index that gives the start and end positions of each molecule in the source SD File. This line index can be used by the downstream `read.SDFindex` function to retrieve specific molecules of interest from the source SD File without reading the entire file into R. The following outlines the typical workflow of this streaming functionality in *ChemmineR*.

Create sample SD File with 100 molecules:

```
> write.SDF(sdfset, "test.sdf")
```

Define descriptor set in a simple function:

```
> desc <- function(sdfset) {
+   cbind(SDFID=sdfid(sdfset),
+         # datablock2ma(datablocklist=datablock(sdfset)),
+         MW=MW(sdfset),
+         groups(sdfset),
+         APFP=desc2fp(x=sdf2ap(sdfset), descnames=1024, type="character"),
+         AP=sdf2ap(sdfset, type="character"),
+         rings(sdfset, type="count", upper=6, arom=TRUE)
+   )
+ }
```

Run `sdfStream` with `desc` function and write results to a file called `matrix.xls`:

```
> sdfStream(input="test.sdf", output="matrix.xls", fct=desc, Nlines=1000)
> # 'Nlines': number of lines to read from input SD File at a time
```

One can also start reading from a specific line number in the SD file. The following example starts at line number 950. This is useful for restarting and debugging the process. With `append=TRUE` the result can be appended to an existing file.

```
> sdfStream(input="test.sdf", output="matrix2.xls", append=FALSE, fct=desc,
+   Nlines=1000, startline=950)
```

Select molecules meeting certain property criteria from SD File using line index generated by previous `sdfStream` step:

```
> indexDF <- read.delim("matrix.xls", row.names=1)[,1:4]
> indexDFsub <- indexDF[indexDF$MW < 400, ]
> # Selects molecules with MW < 400
> sdfset <- read.SDFindex(file="test.sdf", index=indexDFsub, type="SDFset")
> # Collects results in 'SDFset' container
```

Write results directly to SD file without storing larger numbers of molecules in memory:

```
> read.SDFindex(file="test.sdf", index=indexDFsub, type="file", outfile="sub.sdf")
```

Read AP/APFP strings from file into `APset` or `FP` object:

```
> apset <- read.AP(x="matrix.xls", type="ap", colid="AP")
> apfp <- read.AP(x="matrix.xls", type="fp", colid="APFP")
```

Alternatively, one can provide the AP/APFP strings in a named character vector:

```
> apset <- read.AP(x=sdf2ap(sdfset[1:20], type="character"), type="ap")
> fpchar <- desc2fp(sdf2ap(sdfset[1:20]), descnames=1024, type="character")
> fpset <- as(fpchar, "FPset")
```

11 Storing Compounds in an SQL Database

As an alternative to `sdfStream`, there is now also an option to store data in an SQL database, which then allows for fast queries and compound retrieval. This is still an experimental feature. The default database is SQLite, but any other SQL database should work with some minor modifications to the table definitions, which are stored in `schema/compounds.SQLite` under the ChemmineR package directory. Compounds are stored in their entirety in the databases so there is no need to keep any original data files.

Users can define their own set of compound features to compute and store when loading new compounds. Each of these features will be stored in its own, indexed table. Searches can then be performed using these features to quickly find specific compounds. Compounds can always be retrieved quickly because of the database index, no need to scan a large compound file. In addition to user defined features, descriptors can also be computed and stored for each compound.

A new database can be created with the `initDb` function. This takes either an existing database connection, or a filename. If a filename is given then an SQLite database connection is created. It then ensures that the required tables exist and creates them if not. The connection object is then returned. This function can be called safely on the same connection or database many times and will not delete any data.

11.1 Loading Data

The function `loadSdf` can be used to load SDF data, either from a file or *SDFset* object. The `fct` parameter should be a function to extract features from the data. It will be handed an *SDFset* generated from the data being loaded. This may be done in batches, so there is no guarantee that the given *SDFset* will contain the whole dataset. This function should return a data frame with a column for each feature and a row for each compound given. The order of the final data frame should be the same as that of the *SDFset*. The column names will become the feature names. Each of these features will become a new, indexed, table in the database which can be used later to search for compounds.

The `descriptors` parameter can be a function which computes descriptors. This function will also be given an *SDFset* object, which may be done in batches. It should return a data frame with the following two columns: "descriptor" and "descriptor_type". The "descriptor" column should contain a string representation of the descriptor, and "descriptor_type" is the type of the descriptor. Our convention for atom pair is "ap" and "fp" for finger print. The order should also be maintained.

When the data has been loaded, `loadSdf` will return the compound id numbers of each compound loaded. These compound id numbers are computed by the database and are not extracted from the compound data itself. They can be used to quickly retrieve compounds later.

New features can also be added using this function. However, all compounds must have all features so if new features are added to a new set of compounds, all existing features must be computable by the `fct` function given. If new features are detected, all existing compounds will be run through `fct` in order to compute the new features for them as well.

For example, if dataset X is loaded with features F1 and F2, and then at a later time we load dataset Y with new feature F3, the `fct` function used to load dataset Y must compute and return features F1, F2, and F3. `loadSdf` will call `fct` with both datasets X and Y so that all features are available for all compounds. If any features are missing an error will be raised. If just new features are being added, but no new compounds, use the `addNewFeatures` function.

In this example, we create a new database called "test.db" and load it with data from an *SDFset*. We also define `fct` to compute the molecular weight, "MW", and the number of rings and aromatic rings. The `rings` function actually returns a data frame with columns "RINGS" and "AROMATIC", which will be merged into the data frame being created which will also contain the "MW" column. These will be the names used for these features and must be used when searching with them. Finally, the new compound ids are returned and stored in the "ids" variable.

```
> data(sdfsample)
> #create and initialize a new SQLite database
> conn <- initDb("test.db")

[1] "creating db"

> # load data and compute 3 features: molecular weight, with the MW function,
> # and counts for RINGS and AROMATIC, as computed by rings, which
> # returns a data frame itself.
> ids<-loadSdf(conn,sdfsample,
+   function(sdfset)
+     data.frame(MW = MW(sdfset),
+               rings(sdfset,type="count",upper=6, arom=TRUE))
+   )
```

11.2 Searching

Compounds can be searched for using the `findCompounds` function. This function takes a connection object, a vector of feature names used in the tests, and finally, a vector of tests that must all pass for a compound to be included in the result set. Each test should be a boolean expression. For example: "c("MW <= 400","RINGS > 3")" would return all compounds with a molecular weight of 400 or less and more than 3 rings, assuming these features exist in the database. The syntax for each test is "<feature name> <SQL operator> <value>". If you know SQL you can go beyond this basic syntax. These tests will simply be concatenated together with "AND" in-between them and tacked on the end of a WHERE clause of an SQL statement. So any SQL that will work in that context is fine. The function will return a list of compound ids, the actual compounds can be fetched with `getCompounds`. If just the names are needed, the `getCompoundNames` function can be used. Compounds can also be fetched by name using the `findCompoundsByName` function.

In this example we search for compounds with molecular weight less than 300. We then fetch the matching compounds and show their molecular weight.

```
> lightIds <- findCompounds(conn,"MW",c("MW < 300"))
> MW(getCompounds(conn,lightIds))

      206      214      217      222      224      228      229      230      233      234
262.3043 240.2541 265.3050 287.3999 266.7465 278.2789 282.3171 270.3479 223.2252 265.3098
      236      237      242      246      250      255      256      257      268      273
249.2162 285.2581 229.3159 248.3042 266.3427 275.3859 299.8361 263.3752 296.3205 275.3693
      276      282      289      295
294.3510 140.1399 294.3726 276.3357

> #names of matching compounds:
> getCompoundNames(conn,lightIds)

[1] "650006" "650014" "650017" "650023" "650025" "650029" "650030" "650031" "650034" "650035"
[11] "650037" "650038" "650043" "650047" "650052" "650060" "650061" "650062" "650073" "650078"
[21] "650081" "650088" "650095" "650101"
```

12 Working with SDF/SDFset Classes

Several methods are available to return the different data components of *SDF/SDFset* containers in batches. The following examples list the most important ones. To save space their content is not printed in the manual.

```
> view(sdfset[1:4]) # Summary view of several molecules
> length(sdfset) # Returns number of molecules
> sdfset[[1]] # Returns single molecule from SDFset as SDF object
> sdfset[[1]][[2]] # Returns atom block from first compound as matrix
> sdfset[[1]][[2]][1:4,]
> c(sdfset[1:4], sdfset[5:8]) # Concatenation of several SDFsets
```

The `grepSDFset` function allows string matching/searching on the different data components in *SDFset*. By default the function returns a SDF summary of the matching entries. Alternatively, an index of the matches can be returned with the setting `mode="index"`.

```
> grepSDFset("650001", sdfset, field="datablock", mode="subset")
> # To return index, set mode="index"
> .
```

Utilities to maintain unique compound IDs:

```
> sdfid(sdfset[1:4])
> # Retrieves CMP IDs from Molecule Name field in header block.
> cid(sdfset[1:4])
> # Retrieves CMP IDs from ID slot in SDFset.
> unique_ids <- makeUnique(sdfid(sdfset))
> # Creates unique IDs by appending a counter to duplicates.
> cid(sdfset) <- unique_ids # Assigns uniquified IDs to ID slot
```

Subsetting by character, index and logical vectors:

```
> view(sdfset[c("650001", "650012")])
> view(sdfset[4:1])
> mylog <- cid(sdfset) %in% c("650001", "650012")
> view(sdfset[mylog])
```

Accessing *SDF/SDFset* components: header, atom, bond and data blocks:


```

> atomblock(sdf); sdf[[2]]; sdf[["atomblock"]]
> # All three methods return the same component
> header(sdfset[1:4])
> atomblock(sdfset[1:4])
> bondblock(sdfset[1:4])
> datablock(sdfset[1:4])
> header(sdfset[[1]])
> atomblock(sdfset[[1]])
> bondblock(sdfset[[1]])
> datablock(sdfset[[1]])

```

Replacement Methods:

```

> sdfset[[1]][[2]][1,1] <- 999
> atomblock(sdfset)[1] <- atomblock(sdfset)[2]
> datablock(sdfset)[1] <- datablock(sdfset)[2]

```

Assign matrix data to data block:

```

> datablock(sdfset) <- as.matrix(iris[1:100,])
> view(sdfset[1:4])

```

Class coercions from *SDFstr* to *list*, *SDF* and *SDFset*:

```

> as(sdfstr[1:2], "list")
> as(sdfstr[[1]], "SDF")
> as(sdfstr[1:2], "SDFset")

```

Class coercions from *SDF* to *SDFstr*, *SDFset*, *list* with SDF sub-components:

```

> sdfcomplist <- as(sdf, "list")
> sdfcomplist <- as(sdfset[1:4], "list"); as(sdfcomplist[[1]], "SDF")
> sdflist <- as(sdfset[1:4], "SDF"); as(sdflist, "SDFset")
> as(sdfset[[1]], "SDFstr")
> as(sdfset[[1]], "SDFset")

```

Class coercions from *SDFset* to lists with components consisting of SDF or sub-components:

```

> as(sdfset[1:4], "SDF")
> as(sdfset[1:4], "list")
> as(sdfset[1:4], "SDFstr")

```

13 Molecular Property Functions (Physicochemical Descriptors)

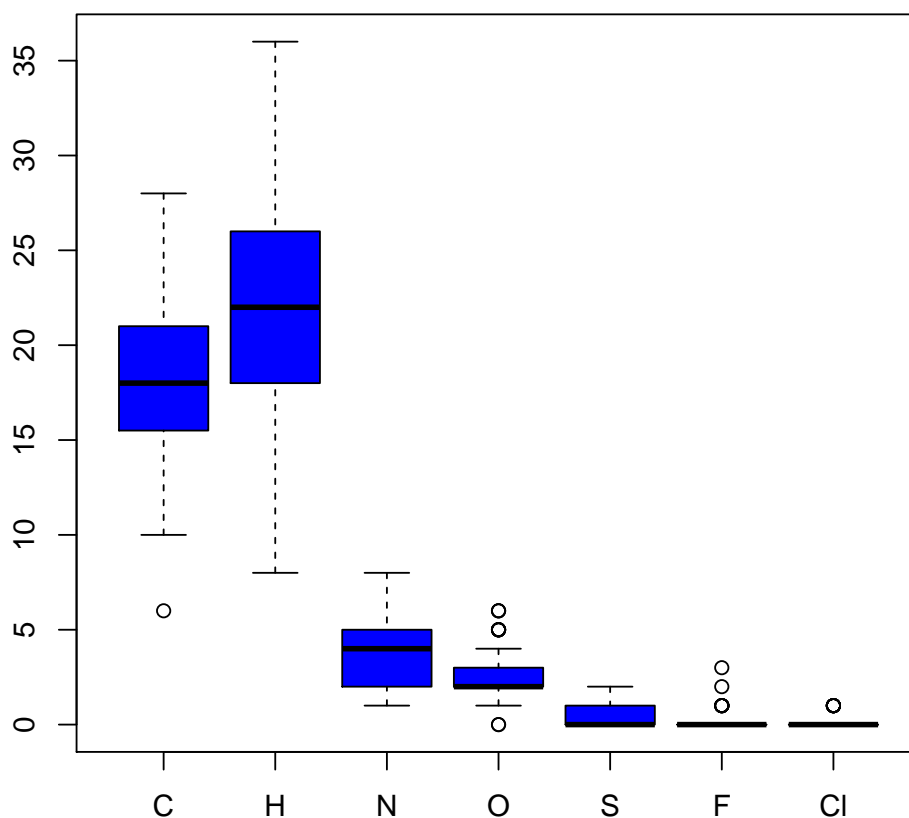
Several methods and functions are available to compute basic compound descriptors, such as molecular formula (MF), molecular weight (MW), and frequencies of atoms and functional groups. In many of these functions, it is important to set `addH=TRUE` in order to include/add hydrogens that are often not specified in an SD file.

```

> propma <- atomcountMA(sdfset, addH=FALSE)
> boxplot(propma, col="blue", main="Atom Frequency")

```

Atom Frequency



```
> boxplot(rowSums(propma), main="All Atom Frequency")
```

Data frame provided by library containing atom names, atom symbols, standard atomic weights, group and period numbers:

```
> data(atomprop)
> atomprop[1:4,]
```

	Number	Name	Symbol	Atomic_weight	Group	Period
1	1	hydrogen	H	1.007940	1	1
2	2	helium	He	4.002602	18	1
3	3	lithium	Li	6.941000	1	2
4	4	beryllium	Be	9.012182	2	2

Compute MW and formula:

```
> MW(sdfset[1:4], addH=FALSE)
```

CMP1	CMP2	CMP3	CMP4
456.4916	357.4069	370.4255	461.5346

```
> MF(sdfset[1:4], addH=FALSE)
```

```

      CMP1      CMP2      CMP3      CMP4
"C23H28N4O6" "C18H23N5O3" "C18H18N4O3S" "C21H27N5O5S"

```

Enumerate functional groups:

```
> groups(sdfset[1:4], groups="fctgroup", type="countMA")
```

	RNH2	R2NH	R3N	ROPO3	ROH	RCHO	RCOR	RCOOH	RCOOR	ROR	RCCH	RCN
CMP1	0	2	1	0	0	0	0	0	0	2	0	0
CMP2	0	2	2	0	1	0	0	0	0	0	0	0
CMP3	0	1	1	0	1	0	1	0	0	0	0	0
CMP4	0	1	3	0	0	0	0	0	0	2	0	0

Combine MW, MF, charges, atom counts, functional group counts and ring counts in one data frame:

```

> propma <- data.frame(MF=MF(sdfset, addH=FALSE), MW=MW(sdfset, addH=FALSE),
+                      Ncharges=sapply(bonds(sdfset, type="charge"), length),
+                      atomcountMA(sdfset, addH=FALSE), groups(sdfset,
+                      type="countMA"), rings(sdfset, upper=6, type="count",
+                      arom=TRUE))
> propma[1:4,]

```

	MF	MW	Ncharges	C	H	N	O	S	F	Cl	RNH2	R2NH	R3N	ROPO3	ROH	RCHO	RCOR	RCOOH
CMP1	C23H28N4O6	456.4916	0	23	28	4	6	0	0	0	0	2	1	0	0	0	0	0
CMP2	C18H23N5O3	357.4069	0	18	23	5	3	0	0	0	0	2	2	0	1	0	0	0
CMP3	C18H18N4O3S	370.4255	0	18	18	4	3	1	0	0	0	1	1	0	1	0	1	0
CMP4	C21H27N5O5S	461.5346	0	21	27	5	5	1	0	0	0	1	3	0	0	0	0	0

	RCOOR	ROR	RCCH	RCN	RINGS	AROMATIC
CMP1	0	2	0	0	4	2
CMP2	0	0	0	0	3	3
CMP3	0	0	0	0	4	2
CMP4	0	2	0	0	3	3

The following shows an example for assigning the values stored in a matrix (e.g. property descriptors) to the data block components in an *SDFset*. Each matrix row will be assigned to the corresponding slot position in the *SDFset*.

```

> datablock(sdfset) <- propma # Works with all SDF components
> datablock(sdfset)[1:4]
> test <- apply(propma[1:4,], 1, function(x) data.frame(col=colnames(propma), value=x))
> sdf.visualize(sdfset[1:4], extra = test)

```

The data blocks in SDFs contain often important annotation information about compounds. The *datablock2ma* function returns this information as matrix for all compounds stored in an *SDFset* container. The *splitNumChar* function can then be used to organize all numeric columns in a *numeric matrix* and the character columns in a *character matrix* as components of a *list* object.

```

> datablocktag(sdfset, tag="PUBCHEM_NIST_INCHI")
> datablocktag(sdfset, tag="PUBCHEM_OPENEYE_CAN_SMILES")

```

Convert entire data block to matrix:

```

> blockmatrix <- datablock2ma(datablocklist=datablock(sdfset))
> # Converts data block to matrix
> numchar <- splitNumChar(blockmatrix=blockmatrix)
> # Splits matrix to numeric matrix and character matrix
> numchar[[1]][1:4,]; numchar[[2]][1:4,]
> # Splits matrix to numeric matrix and character matrix
> .

```

14 Bond Matrices

Bond matrices provide an efficient data structure for many basic computations on small molecules. The function `conMA` creates this data structure from *SDF* and *SDFset* objects. The resulting bond matrix contains the atom labels in the row/column titles and the bond types in the data part. The labels are defined as follows: 0 is no connection, 1 is a single bond, 2 is a double bond and 3 is a triple bond.

```
> conMA(sdfset[1:2], exclude=c("H"))
> # Create bond matrix for first two molecules in sdfset
> conMA(sdfset[[1]], exclude=c("H"))
> # Return bond matrix for first molecule
> plot(sdfset[1], atomnum = TRUE, noHbonds=FALSE, no_print_atoms = "", atomcex=0.8)
> # Plot its structure with atom numbering
> rowSums(conMA(sdfset[[1]], exclude=c("H")))
> # Return number of non-H bonds for each atom
> .
```

15 Charges and Missing Hydrogens

The function `bonds` returns information about the number of bonds, charges and missing hydrogens in *SDF* and *SDFset* objects. It is used by many other functions (e.g. `MW`, `MF`, `atomcount`, `atomcuntMA` and `plot`) to correct for missing hydrogens that are often not specified in SD files.

```
> bonds(sdfset[[1]], type="bonds")[1:4,]
```

	atom	Nbondcount	Nbondrule	charge
1	0	2	2	0
2	0	2	2	0
3	0	2	2	0
4	0	2	2	0

```
> bonds(sdfset[1:2], type="charge")
```

```
$CMP1
NULL
```

```
$CMP2
NULL
```

```
> bonds(sdfset[1:2], type="addNH")
```

```
CMP1 CMP2
0 0
```

16 Ring Perception and Aromaticity Assignment

The function `rings` identifies all possible rings in one or many molecules (here `sdfset[1]`) using the exhaustive ring perception algorithm from Hanser et al. (1996). In addition, the function can return all smallest possible rings as well as aromaticity information.

The following example returns all possible rings in a *list*. The argument `upper` allows to specify an upper length limit for rings. Choosing smaller length limits will reduce the search space resulting in shortened compute times. Note: each ring is represented by a character vector of atom symbols that are numbered by their position in the atom block of the corresponding *SDF*/*SDFset* object.

```
> (ringatoms <- rings(sdfset[1], upper=Inf, type="all", arom=FALSE, inner=FALSE))
```

```
$ring1
```

```
[1] "N_10" "O_6"  "C_32" "C_31" "C_30"
```

 $\$ring2$

```
[1] "C_12" "C_14" "C_15" "C_13" "C_11"
```

 $\$ring3$

```
[1] "C_23" "O_2" "C_27" "C_28" "O_3" "C_25"
```

\$ring4

```
[1] "C_23" "C_21" "C_18" "C_22" "C_26" "C_25"
```

\$ring5

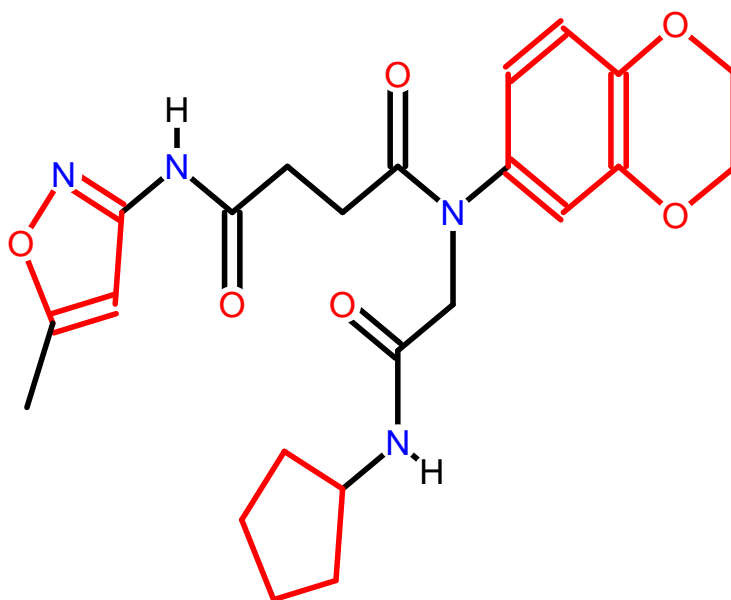
```
[1] "0_3"  "C_28" "C_27" "0_2"  "C_23" "C_21" "C_18" "C_22" "C_26" "C_25"
```

For visual inspection, the corresponding compound structure can be plotted with the ring bonds highlighted in color:

```
> atomindex <- as.numeric(gsub(".*_", "", unique(unlist(ringatoms))))
```

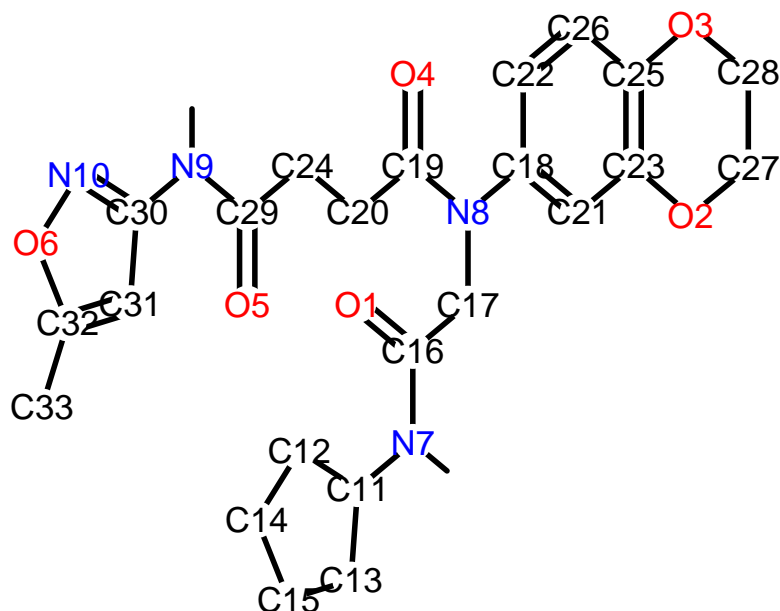
```
> plot(sdfset[1], print=FALSE, colbonds=atomindex)
```

CMP1



Alternatively, one can include the atom numbers in the plot:

```
> plot(sdfset[1], print=FALSE, atomnum=TRUE, no_print_atoms="H")
```

CMP1

Aromaticity information of the rings can be returned in a logical vector by setting `arom=TRUE`:

```
> rings(sdfset[1], upper=Inf, type="all", arom=TRUE, inner=FALSE)
```

```
$RINGS
```

```
$RINGS$ring1
```

```
[1] "N_10" "O_6" "C_32" "C_31" "C_30"
```

```
$RINGS$ring2
```

```
[1] "C_12" "C_14" "C_15" "C_13" "C_11"
```

```
$RINGS$ring3
```

```
[1] "C_23" "O_2" "C_27" "C_28" "O_3" "C_25"
```

```
$RINGS$ring4
```

```
[1] "C_23" "C_21" "C_18" "C_22" "C_26" "C_25"
```

```
$RINGS$ring5
```

```
[1] "O_3" "C_28" "C_27" "O_2" "C_23" "C_21" "C_18" "C_22" "C_26" "C_25"
```

```
$AROMATIC
ring1 ring2 ring3 ring4 ring5
TRUE FALSE FALSE TRUE FALSE
```

Return rings with no more than 6 atoms that are also aromatic:

```
> rings(sdfset[1], upper=6, type="arom", arom=TRUE, inner=FALSE)
```

```
$AROMATIC_RINGS
$AROMATIC_RINGS$ring1
[1] "N_10" "O_6" "C_32" "C_31" "C_30"
```

```
$AROMATIC_RINGS$ring4
[1] "C_23" "C_21" "C_18" "C_22" "C_26" "C_25"
```

Count shortest possible rings and their aromaticity assignments by setting `type=count` and `inner=TRUE`. The inner (smallest possible) rings are identified by first computing all possible rings and then selecting only the inner rings. For more details, consult the help documentation with `?rings`.

```
> rings(sdfset[1:4], upper=Inf, type="count", arom=TRUE, inner=TRUE)
```

	RINGS	AROMATIC
CMP1	4	2
CMP2	3	3
CMP3	4	2
CMP4	3	3

17 Rendering Chemical Structure Images

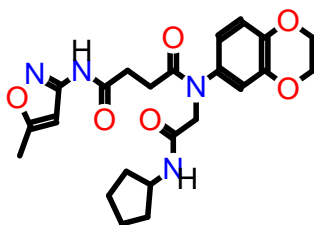
17.1 R Graphics Device

A new plotting function for compound structures has been added to the package recently. This function uses the native R graphics device for generating compound depictions. At this point this function is still in an experimental developmental stage but should become stable soon.

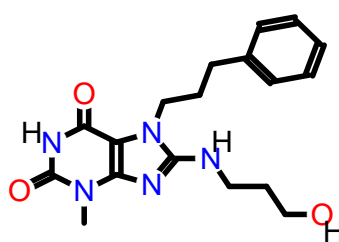
Plot compound Structures with R's graphics device:

```
> data(sdfsample)
> sdfset <- sdfsample
> plot(sdfset[1:4], print=FALSE) # 'print=TRUE' returns SDF summaries
```

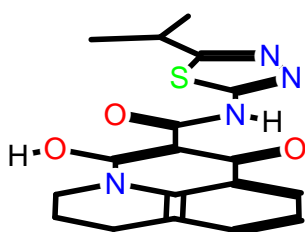
CMP1



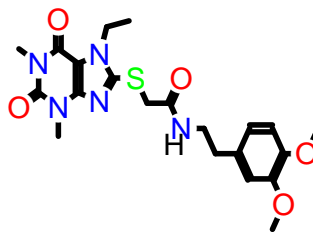
CMP2



CMP3



CMP4

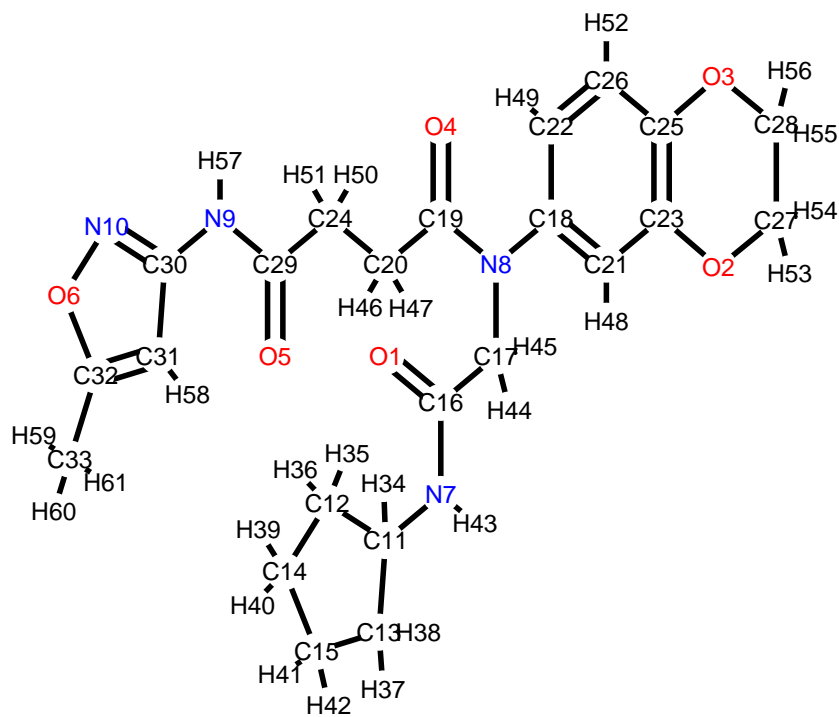


Customized plots:

```
> plot(sdfset[1:4], griddim=c(2,2), print_cid=letters[1:4], print=FALSE,
+       noHbonds=FALSE)
```

In the following plot, the atom block position numbers in the SDF are printed next to the atom symbols (`atomnum = TRUE`). For more details, consult help documentation with `?plotStruc` or `?plot`.

```
> plot(sdfset["CMP1"], atomnum = TRUE, noHbonds=F, no_print_atoms = "",
+       atomcex=0.8, sub=paste("MW:", MW(sdfsampl["CMP1"])), print=FALSE)
```

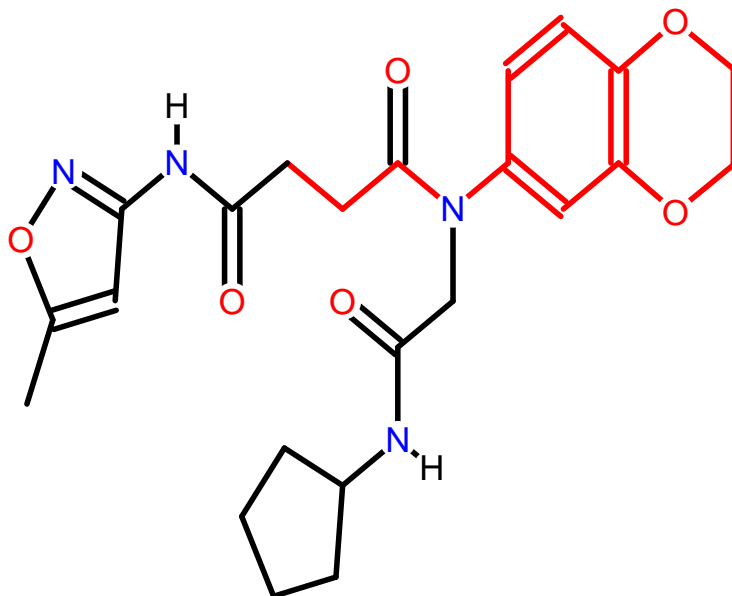

CMP1

MW: 456.49162

Substructure highlighting by atom numbers:

```
> plot(sdfset[1], print=FALSE, colbonds=c(22,26,25,3,28,27,2,23,21,18,8,19,20,24))
```

CMP1



17.2 Online with ChemMine Tools

Alternatively, one can visualize compound structures with a standard web browser using the online ChemMine Tools service. The service allows to display other information next to the structures using the extra argument of the `sdf.visualize` function. The following examples demonstrate, how one can plot and annotate structures by passing on extra data as vector of character strings, matrices or lists.

Plot structures using web service ChemMine Tools:

```
> sdf.visualize(sdfset[1:4])
```

Add extra annotation as *vector*:

```
> sdf.visualize(sdfset[1:4], extra=month.name[1:4])
```

Add extra annotation as *matrix*:

```
> extra <- apply(propma[1:4,], 1, function(x)
+   data.frame(Property=colnames(propma), Value=x))
> sdf.visualize(sdfset[1:4], extra=extra)
```

Add extra annotation as *list*:

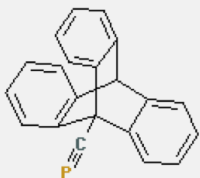
```
> sdf.visualize(sdfset[1:4], extra=bondblock(sdfset[1:4]))
```

[View Previously Accessed Compounds >>>](#)

Width of information table:

Reference Compound (ka-01834)

[View SDF](#)[Structure Search](#)[Add to Selection](#)

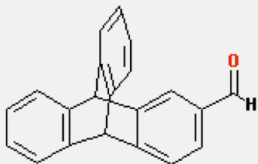


Similarities With All

ids	scores
3	0.550335570
43	0.484662577
42	0.484662577
1	0.484662577
4	0.480122324
2	0.480122324
44	0.356097561
46	0.312500000
11	0.311653117
35	0.287719298

(ChemmineR_Unnamed_Compound_3)

[View SDF](#)[Structure Search](#)[Add to Selection](#)

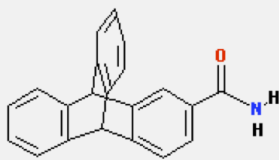


Similarities With All

ids	scores
3	1.000000000
43	0.785977860
42	0.785977860
1	0.785977860
2	0.766423358
4	0.646258503
44	0.561797753
11	0.415204678
35	0.390151515
46	0.368539326

(ChemmineR_Unnamed_Compound_43)

[View SDF](#)[Structure Search](#)[Add to Selection](#)



Similarities With All

ids	scores
43	1.000000000
42	0.840000000
1	0.840000000
3	0.785977860
2	0.709459459
4	0.611464968
44	0.601108033
11	0.390109890
46	0.339702760
35	0.223128252

Figure 3: Visualization webpage created by calling `sdf.visualize`.

27

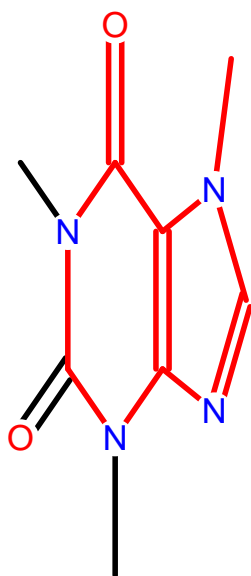
18 Similarity Comparisons and Searching

18.1 Maximum Common Substructure (MCS) Searching

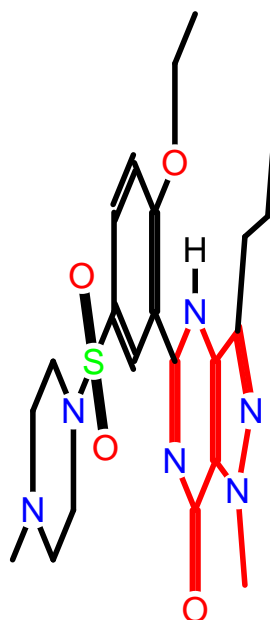
The ChemmineR add-on package `fmcsR` provides support for identifying maximum common substructures (MCSs) and flexible MCSs among compounds. The algorithm can be used for pairwise compound comparisons, structure similarity searching and clustering. The manual describing this functionality is available [here](#) and the associated publication is available in Wang et al. (2013). The following gives a short preview of some functionalities provided by the `fmcsR` package.

```
> library(fmcsR)
> data(fmcstest) # Loads test sdfset object
> test <- fmcs(fmcstest[1], fmcstest[2], au=2, bu=1) # Searches for MCS with mismatches
> plotMCS(test) # Plots both query compounds with MCS in color
```

Caffeine



Viagra



18.2 AP/APset Classes for Storing Atom Pair Descriptors

The function `sdf2ap` computes atom pair descriptors for one or many compounds (Carhart et al., 1985; Chen and Reynolds, 2002). It returns a searchable atom pair database stored in a container of class `APset`, which can be used for structural similarity searching and clustering. As similarity measure, the Tanimoto coefficient or related coefficients can

be used. An *APset* object consists of one or many *AP* entries each storing the atom pairs of a single compound. Note: the deprecated `cmp.parse` function is still available which also generates atom pair descriptor databases, but directly from an SD file. Since the latter function is less flexible it may be discontinued in the future.

Generate atom pair descriptor database for searching:

```
> ap <- sdf2ap(sdfset[[1]]) # For single compound
> ap
```

An instance of "AP"

```
<<atom pairs>>
```

```
52614450304 52615497856 52615514112 52616547456 52616554624 ... length: 528
```

```
> apset <- sdf2ap(sdfset) # For many compounds.
```

```
> view(apset[1:4])
```

```
$`650001`
```

An instance of "AP"

```
<<atom pairs>>
```

```
53688190976 53688190977 53688190978 53688190979 53688190980 ... length: 528
```

```
$`650002`
```

An instance of "AP"

```
<<atom pairs>>
```

```
53688190976 53688190977 53688190978 53688190979 53689239552 ... length: 325
```

```
$`650003`
```

An instance of "AP"

```
<<atom pairs>>
```

```
52615496704 53688190976 53688190977 53689239552 53697627136 ... length: 325
```

```
$`650004`
```

An instance of "AP"

```
<<atom pairs>>
```

```
52617593856 52618642432 52619691008 52619691009 52628079616 ... length: 496
```

Return main components of *APset* objects:

```
> cid(apset[1:4]) # Compound IDs
> ap(apset[1:4]) # Atom pair descriptors
> db.explain(apset[1]) # Return atom pairs in human readable format
```

Coerce *APset* to other objects:

```
> apset2descdb(apset) # Returns old list-style AP database
> tmp <- as(apset, "list") # Returns list
> as(tmp, "APset") # Converts list back to APset
```

18.3 Large SDF and Atom Pair Databases

When working with large data sets it is often desirable to save the *SDFset* and *APset* containers as binary R objects to files for later use. This way they can be loaded very quickly into a new R session without recreating them every time from scratch.

Save and load of *SDFset* and *APset* containers:

```
> save(sdfset, file = "sdfset.rda", compress = TRUE)
> load("sdfset.rda")
> save(apset, file = "apset.rda", compress = TRUE)
> load("apset.rda")
```

18.4 Pairwise Compound Comparisons with Atom Pairs

The `cmp.similarity` function computes the atom pair similarity between two compounds using the Tanimoto coefficient as similarity measure. The coefficient is defined as $c/(a+b+c)$, which is the proportion of the atom pairs shared among two compounds divided by their union. The variable c is the number of atom pairs common in both compounds, while a and b are the numbers of their unique atom pairs.

```
> cmp.similarity(apset[1], apset[2])

[1] 0.2637037

> cmp.similarity(apset[1], apset[1])

[1] 1
```

18.5 Similarity Searching with Atom Pairs

The `cmp.search` function searches an atom pair database for compounds that are similar to a query compound. The following example returns a data frame where the rows are sorted by the Tanimoto similarity score (best to worst). The first column contains the indices of the matching compounds in the database. The argument `cutoff` can be a similarity cutoff, meaning only compounds with a similarity value larger than this cutoff will be returned; or it can be an integer value restricting how many compounds will be returned. When supplying a cutoff of 0, the function will return the similarity values for every compound in the database.

```
> cmp.search(apset, apset["650065"], type=3, cutoff = 0.3, quiet=TRUE)

  index  cid  scores
1    61 650066 1.0000000
2    60 650065 1.0000000
3    67 650072 0.3389831
4    11 650011 0.3190608
5    15 650015 0.3184524
6    86 650092 0.3154270
7    64 650069 0.3010279
```

Alternatively, the function can return the matches in form of an index or a named vector if the `type` argument is set to 1 or 2, respectively.

```
> cmp.search(apset, apset["650065"], type=1, cutoff = 0.3, quiet=TRUE)

[1] 61 60 67 11 15 86 64

> cmp.search(apset, apset["650065"], type=2, cutoff = 0.3, quiet=TRUE)

  650066  650065  650072  650011  650015  650092  650069
1.0000000 1.0000000 0.3389831 0.3190608 0.3184524 0.3154270 0.3010279
```

18.6 FP/FPset Classes for Storing Fingerprints

The *FPset* class stores fingerprints of small molecules in a matrix-like representation where every molecule is encoded as a fingerprint of the same type and length. The *FPset* container acts as a searchable database that contains the fingerprints of many molecules. The *FP* container holds only one fingerprint. Several constructor and coerce methods are provided to populate *FP/FPset* containers with fingerprints, while supporting any type and length of fingerprints. For instance, the function `desc2fp` generates fingerprints from an atom pair database stored in an *APset*, and `as(matrix, "FPset")` and `as(character, "FPset")` construct an *FPset* database from objects where the fingerprints are represented as *matrix* or *character* objects, respectively.

Show slots of *FPset* class:

```
> showClass("FPset")
```

```
Class "FPset" [package "ChemmineR"]
```

Slots:

```
Name:    fpma
```

```
Class: matrix
```

Instance of *FPset* class:

```
> data(apset)
```

```
> (fpset <- desc2fp(apset))
```

An instance of a 1024 bit "FPset" with 100 molecules

```
> view(fpset[1:2])
```

```
$`650001`
```

An instance of "FP"

```
<<fingerprint>>
```

```
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 ... length: 1024
```

```
$`650002`
```

An instance of "FP"

```
<<fingerprint>>
```

```
1 1 1 1 1 1 1 1 1 0 1 1 1 1 0 0 0 1 1 ... length: 1024
```

FPset class usage:

```
> fpset[1:4] # behaves like a list
```

An instance of a 1024 bit "FPset" with 4 molecules

```
> fpset[[1]] # returns FP object
```

An instance of "FP"

```
<<fingerprint>>
```

```
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 ... length: 1024
```

```
> length(fpset) # number of compounds
```

```
[1] 100
```

```
> cid(fpset) # returns compound ids
```

```
[1] "650001" "650002" "650003" "650004" "650005" "650006" "650007" "650008" "650009" "650010"
[11] "650011" "650012" "650013" "650014" "650015" "650016" "650017" "650019" "650020" "650021"
[21] "650022" "650023" "650024" "650025" "650026" "650027" "650028" "650029" "650030" "650031"
[31] "650032" "650033" "650034" "650035" "650036" "650037" "650038" "650039" "650040" "650041"
[41] "650042" "650043" "650044" "650045" "650046" "650047" "650048" "650049" "650050" "650052"
[51] "650054" "650056" "650058" "650059" "650060" "650061" "650062" "650063" "650064" "650065"
[61] "650066" "650067" "650068" "650069" "650070" "650071" "650072" "650073" "650074" "650075"
[71] "650076" "650077" "650078" "650079" "650080" "650081" "650082" "650083" "650085" "650086"
[81] "650087" "650088" "650089" "650090" "650091" "650092" "650093" "650094" "650095" "650096"
[91] "650097" "650098" "650099" "650100" "650101" "650102" "650103" "650104" "650105" "650106"
```

```
> fpset[10] <- 0 # replacement of 10th fingerprint to all zeros
> cid(fpset) <- 1:length(fpset) # replaces compound ids
> c(fpset[1:4], fpset[11:14]) # concatenation of several FPset objects
```

An instance of a 1024 bit "FPset" with 8 molecules

Construct *FPset* class from *matrix*:

```
> fpma <- as.matrix(fpset) # coerces FPset to matrix
> as(fpma, "FPset")
```

An instance of a 1024 bit "FPset" with 100 molecules

Construct *FPset* class from *character vector*:

```
> fpchar <- as.character(fpset) # coerces FPset to character strings
> as(fpchar, "FPset") # construction of FPset class from character vector
```

An instance of a 1024 bit "FPset" with 100 molecules

Compound similarity searching with *FPset*:

```
> fpSim(fpset[1], fpset, method="Tanimoto", cutoff=0.4, top=4)
```

```
      1      96      67      15
1.0000000 0.4719101 0.4288499 0.4275229
```

18.7 Atom Pair Fingerprints

Atom pairs can be converted into binary atom pair fingerprints of fixed length. Computations on this compact data structure are more time and memory efficient than on their relatively complex atom pair counterparts. The function `desc2fp` generates fingerprints from descriptor vectors of variable length such as atom pairs stored in *APset* or *list* containers. The obtained fingerprints can be used for structure similarity comparisons, searching and clustering.

Create atom pair sample data set:

```
> data(sdfsampl)
> sdfset <- sdfsampl[1:10]
> apset <- sdf2ap(sdfset)
```

Compute atom pair fingerprint database using internal atom pair selection containing the 4096 most common atom pairs identified in DrugBank's compound collection. For details see `?apfp`. The following example uses from this set the 1024 most frequent atom pairs:

```
> fpset <- desc2fp(apset, descnames=1024, type="FPset")
```

Alternatively, one can provide any custom atom pair selection. Here, the 1024 most common ones in *apset*:


```
> fpset1024 <- names(rev(sort(table(unlist(as(apset, "list"))))) [1:1024]))
> fpset <- desc2fp(apset, descnames=fpset1024, type="FPset")
```

A more compact way of storing fingerprints is as character values:

```
> fpchar <- desc2fp(x=apset, descnames=1024, type="character")
> fpchar <- as.character(fpset)
```

Converting a fingerprint database to a matrix and vice versa:

```
> fpma <- as.matrix(fpset)
> fpset <- as(fpma, "FPset")
```

Similarity searching and returning Tanimoto similarity coefficients:

```
> fpSim(fpset[1], fpset, method="Tanimoto")
```

Under method one can choose from several predefined similarity measures including Tanimoto (default), Euclidean, Tversky or Dice. Alternatively, one can pass on custom similarity functions.

```
> fpSim(fpset[1], fpset, method="Tversky", cutoff=0.4, top=4, alpha=0.5, beta=1)
```

Example for using a custom similarity function:

```
> myfct <- function(a, b, c, d) c/(a+b+c+d)
> fpSim(fpset[1], fpset, method=myfct)
```

Clustering example:

```
> simMAap <- sapply(cid(apfpset), function(x) fpSim(x=apfpset[x], apfpset, sorted=FALSE))
> hc <- hclust(as.dist(1-simMAap), method="single")
> plot(as.dendrogram(hc), edgePar=list(col=4, lwd=2), horiz=TRUE)
```

18.8 Pairwise Compound Comparisons with PubChem Fingerprints

The `fpSim` function computes the similarity coefficients (e.g. Tanimoto) for pairwise comparisons of binary fingerprints. For this data type, *c* is the number of "on-bits" common in both compounds, and *a* and *b* are the numbers of their unique "on-bits". Currently, the PubChem fingerprints need to be provided (here PubChem's SD files) and cannot be computed from scratch in *ChemmineR*. The PubChem fingerprint specifications can be loaded with `data(pubchemFPencoding)`.

Convert base 64 encoded PubChem fingerprints to *character* vector, *matrix* or *FPset* object:

```
> cid(sdfset) <- sdfid(sdfset)
> fpset <- fp2bit(sdfset, type=1)
> fpset <- fp2bit(sdfset, type=2)
> fpset <- fp2bit(sdfset, type=3)
> fpset
```

An instance of a 881 bit "FPset" with 100 molecules

Pairwise compound structure comparisons:

```
> fpSim(fpset[1], fpset[2])

650002
0.5364807
```

18.9 Similarity Searching with PubChem Fingerprints

Similarly, the `fpSim` function provides search functionality for PubChem fingerprints:

```
> fpSim(fpset["650065"], fpset, method="Tanimoto", cutoff=0.6, top=6)

650065    650066    650035    650019    650012    650046
1.0000000 0.9944751 0.7435897 0.7432432 0.7230047 0.7142857
```

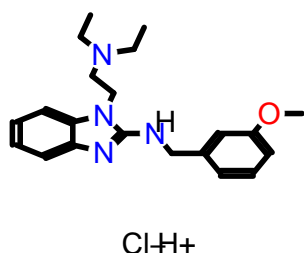
18.10 Visualize Similarity Search Results

The `cmp.search` function allows to visualize the chemical structures for the search results. Similar but more flexible chemical structure rendering functions are `plot` and `sdf.visualize` described above. By setting the `visualize` argument in `cmp.search` to `TRUE`, the matching compounds and their scores can be visualized with a standard web browser. Depending on the `visualize.browse` argument, an URL will be printed or a webpage will be opened showing the structures of the matching compounds along with their scores.

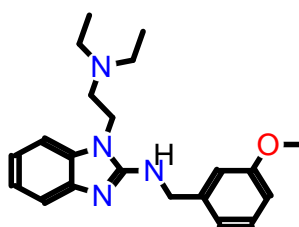
View similarity search results in R's graphics device:

```
> cid(sdfset) <- cid(apset) # Assure compound name consistency among objects.  
> plot(sdfset[names(cmp.search(apset, apset["650065"], type=2, cutoff=4,  
+      quiet=TRUE))], print=FALSE)
```

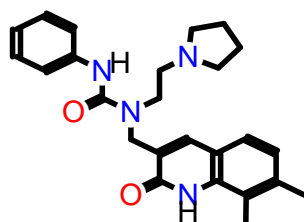
650065



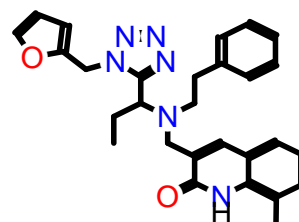
650066



650072



650011



View results online with Chemmine Tools:

```
> similarities <- cmp.search(apset, apset[1], type=3, cutoff = 10)  
> sdf.visualize(sdfset[similarities[,1]], extra=similarities[,3])
```

19 Clustering

19.1 Clustering Identical or Very Similar Compounds

Often it is of interest to identify very similar or identical compounds in a compound set. The `cmp.duplicated` function can be used to quickly identify very similar compounds in atom pair sets, which will be frequently, but not necessarily, identical compounds.

Identify compounds with identical AP sets:

```
> cmp.duplicated(apset, type=1)[1:4] # Returns AP duplicates as logical vector
```

```
[1] FALSE FALSE FALSE FALSE
```

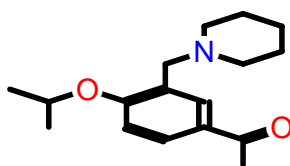
```
> cmp.duplicated(apset, type=2)[1:4,] # Returns AP duplicates as data frame
```

	ids	CLSZ_100	CLID_100
1	650082	1	1
2	650059	2	2
3	650060	2	2
4	650010	1	3

Plot the structure of two pairs of duplicates:

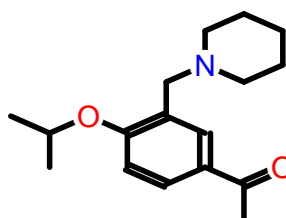
```
> plot(sdfset[c("650059", "650060", "650065", "650066")], print=FALSE)
```

650059

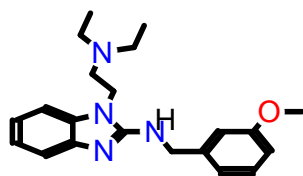


Cl-H+

650060

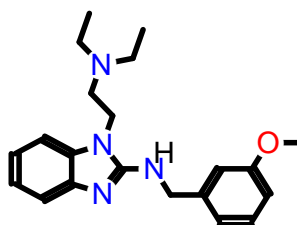


650065



Cl-H+

650066



Remove AP duplicates from SDFset and APset objects:

```
> apdups <- cmp.duplicated(apset, type=1)
> sdfset[which(!apdups)]; apset[which(!apdups)]
```

An instance of "SDFset" with 96 molecules

An instance of "APset" with 96 molecules

Alternatively, one can identify duplicates via other descriptor types if they are provided in the data block of an imported SD file. For instance, one can use here fingerprints, InChI, SMILES or other molecular representations. The following examples show how to enumerate by identical InChI strings, SMILES strings and molecular formula, respectively.

```
> count <- table(datablocktag(sdfset, tag="PUBCHEM_NIST_INCHI"))
> count <- table(datablocktag(sdfset, tag="PUBCHEM_OPENEYE_CAN_SMILES"))
> count <- table(datablocktag(sdfset, tag="PUBCHEM_MOLECULAR_FORMULA"))
> count[1:4]
```

C10H9FN2O2S	C11H12N4OS	C11H13NO4	C12H11ClN2OS
1	1	1	1

19.2 Binning Clustering

Compound libraries can be clustered into discrete similarity groups with the binning clustering function `cmp.cluster`. The function accepts as input an atom pair (APset) or a fingerprint (FPset) descriptor database as well as a similarity threshold. The binning clustering result is returned in form of a data frame. Single linkage is used for cluster joining. The function calculates the required compound-to-compound distance information on the fly, while a memory-intensive distance matrix is only created upon user request via the `save.distances` argument (see below).

Because an optimum similarity threshold is often not known, the `cmp.cluster` function can calculate cluster results for multiple cutoffs in one step with almost the same speed as for a single cutoff. This can be achieved by providing several cutoffs under the `cutoff` argument. The clustering results for the different cutoffs will be stored in one data frame.

One may force the `cmp.cluster` function to calculate and store the distance matrix by supplying a file name to the `save.distances` argument. The generated distance matrix can be loaded and passed on to many other clustering methods available in R, such as the hierarchical clustering function `hclust` (see below).

If a distance matrix is available, it may also be supplied to `cmp.cluster` via the `use.distances` argument. This is useful when one has a pre-computed distance matrix either from a previous call to `cmp.cluster` or from other distance calculation subroutines.

Single-linkage binning clustering with one or multiple cutoffs:

```
> clusters <- cmp.cluster(db=apset, cutoff = c(0.7, 0.8, 0.9), quiet = TRUE)
```

sorting result...

```
> clusters[1:12,]
```

	ids	CLSZ_0.7	CLID_0.7	CLSZ_0.8	CLID_0.8	CLSZ_0.9	CLID_0.9
48	650049	2	48	2	48	2	48
49	650050	2	48	2	48	2	48
54	650059	2	54	2	54	2	54
55	650060	2	54	2	54	2	54
56	650061	2	56	2	56	2	56
57	650062	2	56	2	56	2	56
58	650063	2	58	2	58	2	58
59	650064	2	58	2	58	2	58
60	650065	2	60	2	60	2	60
61	650066	2	60	2	60	2	60
1	650001	1	1	1	1	1	1
2	650002	1	2	1	2	1	2

Clustering of FPset objects with multiple cutoffs. This method allows to call various similarity methods provided by the fpSim function. For details consult ?fpSim.

```
> fpset <- desc2fp(apset)
> clusters2 <- cmp.cluster(fpset, cutoff=c(0.5, 0.7, 0.9), method="Tanimoto",
+                           quiet=TRUE)
```

sorting result...

```
> clusters2[1:12,]

      ids CLSZ_0.5 CLID_0.5 CLSZ_0.7 CLID_0.7 CLSZ_0.9 CLID_0.9
69 650074      14      11        2      69      1      69
79 650085      14      11        2      69      1      79
11 650011      14      11        1     11      1      11
15 650015      14      11        1     15      1      15
45 650046      14      11        1     45      1      45
47 650048      14      11        1     47      1      47
51 650054      14      11        1     51      1      51
53 650058      14      11        1     53      1      53
64 650069      14      11        1     64      1      64
65 650070      14      11        1     65      1      65
67 650072      14      11        1     67      1      67
86 650092      14      11        1     86      1      86
```

Sames as above, but using Tversky similarity measure:

```
> clusters3 <- cmp.cluster(fpset, cutoff=c(0.5, 0.7, 0.9), method="Tversky",
+                           alpha=0.3, beta=0.7, quiet=TRUE)
```

sorting result...

Return cluster size distributions for each cutoff:

```
> cluster.sizestat(clusters, cluster.result=1)
```

```
cluster size count
1          1     90
2          2      5
```

```
> cluster.sizestat(clusters, cluster.result=2)
```

```
cluster size count
1          1     90
2          2      5
```

```
> cluster.sizestat(clusters, cluster.result=3)
```

```
cluster size count
1          1     90
2          2      5
```

Enforce calculation of distance matrix:

```
> clusters <- cmp.cluster(db=apset, cutoff = c(0.65, 0.5, 0.3),
+                           save.distances="distmat.rda")
> # Saves distance matrix to file "distmat.rda" in current working directory.
> load("distmat.rda") # Loads distance matrix.
```

19.3 Jarvis-Patrick Clustering

The Jarvis-Patrick clustering algorithm is widely used in cheminformatics (Jarvis and Patrick, 1973). It requires a nearest neighbor table, which consists of j nearest neighbors for each item (e.g. compound). The nearest neighbor table is then used to join items into clusters when they meet the following requirements: (a) they are contained in each other's neighbor list and (b) they share at least k nearest neighbors. The values for j and k are user-defined parameters. The `jarvisPatrick` function implemented in *ChemmineR* takes a nearest neighbor table generated by `nearestNeighbors`, which works for *APset* and *FPset* objects. This function takes either the standard Jarvis-Patrick j parameter (as the `numNbrs` parameter), or else a cutoff value, which is an extension to the basic algorithm that we have added. Given a cutoff value, the nearest neighbor table returned contains every neighbor with a similarity greater than the cutoff value, for each item. This allows one to generate tighter clusters and to minimize certain limitations of this method, such as false joins of completely unrelated items when operating on small data sets. The `trimNeighbors` function can also be used to take an existing nearest neighbor table and remove all neighbors whose similarity value is below a given cutoff value. This allows one to compute a very relaxed nearest neighbor table initially, and then quickly try different refinements later.

In case an existing nearest neighbor matrix needs to be used, the `fromNNMatrix` function can be used to transform it into the list structure that `jarvisPatrick` requires. The input matrix must have a row for each compound, and each row should be the index values of the neighbors of compound represented by that row. The names of each compound can also be given through the `names` argument. If not given, it will attempt to use the `rownames` of the given matrix.

The `jarvisPatrick` function also allows one to relax some of the requirements of the algorithm through the `mode` parameter. When set to "a1a2b", then all requirements are used. If set to "a1b", then (a) is relaxed to a unidirectional requirement. Lastly, if `mode` is set to "b", then only requirement (b) is used, which means that all pairs of items will be checked to see if (b) is satisfied between them. The size of the clusters generated by the different methods increases in this order: "a1a2b" < "a1b" < "b". The run time of method "a1a2b" follows a close to linear relationship, while it is nearly quadratic for the much more exhaustive method "b". Only methods "a1a2b" and "a1b" are suitable for clustering very large data sets (e.g. >50,000 items) in a reasonable amount of time.

An additional extension to the algorithm is the ability to set the linkage mode. The `linkage` parameter can be one of "single", "average", or "complete", for single linkage, average linkage and complete linkage merge requirements, respectively. In the context of Jarvis-Patrick, average linkage means that at least half of the pairs between the clusters under consideration must meet requirement (b). Similarly, for complete linkage, all pairs must requirement (b). Single linkage is the normal case for Jarvis-Patrick and just means that at least one pair must meet requirement (b).

The output is a cluster vector with the item labels in the name slot and the cluster IDs in the data slot. There is a utility function called `byCluster`, which takes out cluster vector output by `jarvisPatrick` and transforms it into a list of vectors. Each slot of the list is named with a cluster id and the vector contains the cluster members. By default the function excludes singletons from the output, but they can be included by setting `excludeSingletons=FALSE`.

Load/create sample *APset* and *FPset*:

```
> data(apset)
> fpset <- desc2fp(apset)
```

Standard Jarvis-Patrick clustering on *APset* and *FPset* objects:

```
> jarvisPatrick(nearestNeighbors(apset,numNbrs=6), k=5, mode="a1a2b") #Using "APset"
```

650001	650002	650003	650004	650005	650006	650007	650008	650009	650010	650011	650012	650013
1	2	3	4	5	6	7	8	9	10	11	12	13
650014	650015	650016	650017	650019	650020	650021	650022	650023	650024	650025	650026	650027
14	11	15	16	17	18	19	20	21	22	23	24	25
650028	650029	650030	650031	650032	650033	650034	650035	650036	650037	650038	650039	650040
26	27	28	29	30	31	32	33	34	35	36	37	38
650041	650042	650043	650044	650045	650046	650047	650048	650049	650050	650052	650054	650056
39	40	41	42	43	44	45	46	47	48	49	50	51
650058	650059	650060	650061	650062	650063	650064	650065	650066	650067	650068	650069	650070
52	53	54	55	56	57	58	59	60	61	62	63	64
650071	650072	650073	650074	650075	650076	650077	650078	650079	650080	650081	650082	650083
65	66	67	68	69	70	71	72	73	74	75	76	77

```

650085 650086 650087 650088 650089 650090 650091 650092 650093 650094 650095 650096 650097
   78    79    80    81    82    83    84    85    86    87    88    89    90
650098 650099 650100 650101 650102 650103 650104 650105 650106
   91    92    93    94    95    96    97    98    99

```

```
> jarvisPatrick(nearestNeighbors(fpset,numNbrs=6), k=5, mode="a1a2b") #Using "FPset"
```

```

650001 650002 650003 650004 650005 650006 650007 650008 650009 650010 650011 650012 650013
   1     2     3     4     5     6     7     8     9    10    11    12    13
650014 650015 650016 650017 650019 650020 650021 650022 650023 650024 650025 650026 650027
   14    11    15    16    17    18    19    20    21    22    23    24    25
650028 650029 650030 650031 650032 650033 650034 650035 650036 650037 650038 650039 650040
   26    27    28    29    30    31    32    33    34    35    36    37    38
650041 650042 650043 650044 650045 650046 650047 650048 650049 650050 650052 650054 650056
   39    40    41    42    43    44    45    46    47    48    49    50    51
650058 650059 650060 650061 650062 650063 650064 650065 650066 650067 650068 650069 650070
   52    53    54    55    56    57    58    59    60    61    62    63    64
650071 650072 650073 650074 650075 650076 650077 650078 650079 650080 650081 650082 650083
   65    66    67    68    69    70    71    72    73    74    75    76    77
650085 650086 650087 650088 650089 650090 650091 650092 650093 650094 650095 650096 650097
   78    79    80    81    82    83    84    85    86    87    88    89    90
650098 650099 650100 650101 650102 650103 650104 650105 650106
   91    92    93    94     1    95    96    97    98

```

The following example runs Jarvis-Patrick clustering with a minimum similarity cutoff value (here Tanimoto coefficient). In addition, it uses the much more exhaustive "b" method that generates larger cluster sizes, but significantly increased the run time. For more details, consult the corresponding help file with `?jarvisPatrick`.

```
> cl<-jarvisPatrick(nearestNeighbors(fpset,cutoff=0.6, method="Tanimoto"), k=2 ,mode="b")
> byCluster(cl)
```

```
$`11`
[1] "650011" "650092"
```

```
$`15`
[1] "650015" "650069"
```

```
$`45`
[1] "650046" "650054"
```

```
$`48`
[1] "650049" "650050"
```

```
$`52`
[1] "650059" "650060"
```

```
$`53`
[1] "650061" "650062"
```

```
$`54`
[1] "650063" "650064"
```

```
$`55`
[1] "650065" "650066"
```

```
$`62`
[1] "650074" "650085"
```

Output nearest neighbor table (*matrix*):

```
> nnm <- nearestNeighbors(fpset,numNbrs=6)
> nnm$names[1:4]

[1] "650001" "650002" "650003" "650004"

> nnm$ids[1:4,]

NULL

> nnm$similarities[1:4,]

      650001    650102    650072    650015    650094    650069
sim      1 0.4719101 0.4288499 0.4275229 0.4247423 0.4187380
sim      1 0.4343891 0.4246575 0.4216867 0.3939394 0.3922078
sim      1 0.4152249 0.3619303 0.3610315 0.3424242 0.3367089
sim      1 0.5791045 0.4973958 0.4192708 0.4166667 0.4104683
```

Trim nearest neighbor table:

```
> nnm <- trimNeighbors(nnm,cutoff=0.4)
> nnm$similarities[1:4,]

      650001    650102    650072    650015    650094    650069
sim      1 0.4719101 0.4288499 0.4275229 0.4247423 0.4187380
sim      1 0.4343891 0.4246575 0.4216867      NA      NA
sim      1 0.4152249      NA      NA      NA      NA
sim      1 0.5791045 0.4973958 0.4192708 0.4166667 0.4104683

>
```

Perform clustering on precomputed nearest neighbor table:

```
> jarvisPatrick(nnm, k=5,mode="b")

650001 650002 650003 650004 650005 650006 650007 650008 650009 650010 650011 650012 650013
      1      2      3      4      5      6      7      8      9     10     11     12     13
650014 650015 650016 650017 650019 650020 650021 650022 650023 650024 650025 650026 650027
     14     11     15     16     17     18     19     20     21     22     23     24     25
650028 650029 650030 650031 650032 650033 650034 650035 650036 650037 650038 650039 650040
     26     27     28     29     30     31     32     33     34     35     36     37     38
650041 650042 650043 650044 650045 650046 650047 650048 650049 650050 650052 650054 650056
     39     40     41     42     43     11     44     11     45     46     47     48     49
650058 650059 650060 650061 650062 650063 650064 650065 650066 650067 650068 650069 650070
     50     51     52     53     54     55     56     57     57     58     59     11     60
650071 650072 650073 650074 650075 650076 650077 650078 650079 650080 650081 650082 650083
     61     62     63     64     65     66     67     68     69     37     70     71     72
650085 650086 650087 650088 650089 650090 650091 650092 650093 650094 650095 650096 650097
     64     73     74     75     76     77     78     11     79     80     81     82     83
650098 650099 650100 650101 650102 650103 650104 650105 650106
     84     85     86     87      1     88     89     90     91
```

Using a user defined nearest neighbor matrix:

```
> nn <- matrix(c(1,2,2,1),2,2,dimnames=list(c('one','two'))))
> nn
```



```

      [,1] [,2]
one      1    2
two      2    1

> byCluster(jarvisPatrick(fromNNMatrix(nn),k=1))

$`1`
[1] "one" "two"

```

19.4 Multi-Dimensional Scaling (MDS)

To visualize and compare clustering results, the `cluster.visualize` function can be used. The function performs Multi-Dimensional Scaling (MDS) and visualizes the results in form of a scatter plot. It requires as input an *APset*, a clustering result from `cmp.cluster`, and a cutoff for the minimum cluster size to consider in the plot. To help determining a proper cutoff size, the `cluster.sizestat` function is provided to generate cluster size statistics.

MDS clustering and scatter plot:

```

> cluster.visualize(apset, clusters, size.cutoff=2, quiet = TRUE)
> # Color codes clusters with at least two members.
> cluster.visualize(apset, clusters, quiet = TRUE) # Plots all items.

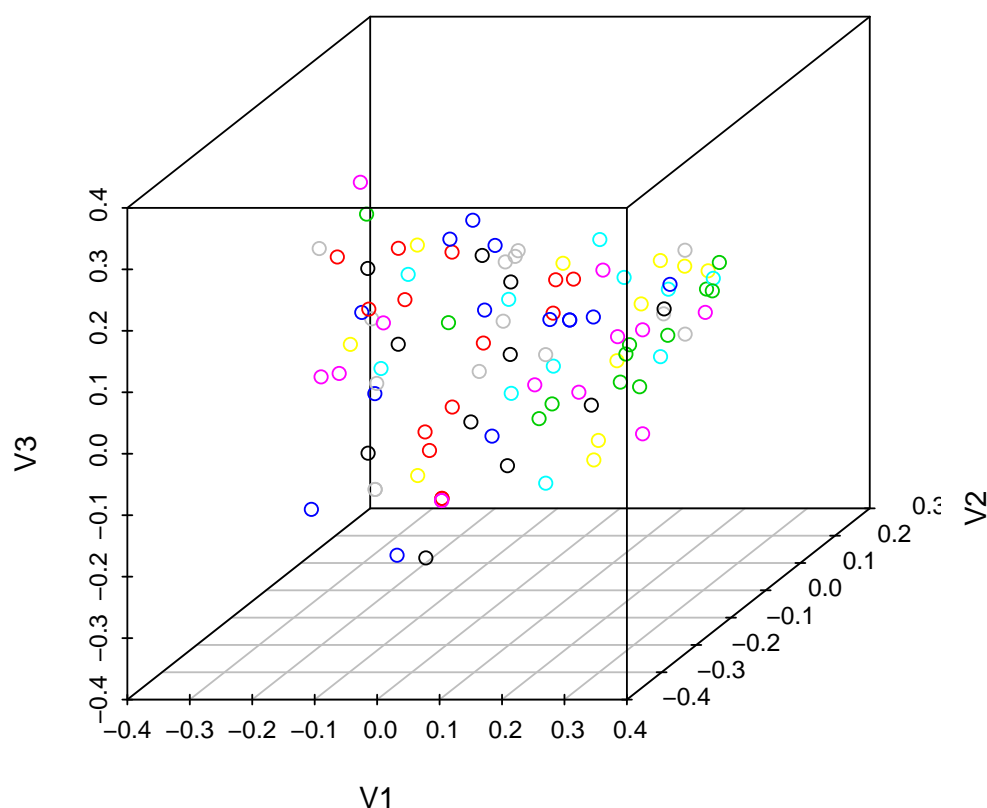
```

Create a 3D scatter plot of MDS result:

```

> library(scatterplot3d)
> coord <- cluster.visualize(apset, clusters, size.cutoff=1, dimensions=3, quiet=TRUE)
> scatterplot3d(coord)

```



Interactive 3D scatter plot with Open GL (graphics not evaluated here):

```
> library(rgl)
> rgl.open(); offset <- 50; par3d(windowRect=c(offset, offset, 640+offset, 640+offset))
> rm(offset); rgl.clear(); rgl.viewpoint(theta=45, phi=30, fov=60, zoom=1)
> spheres3d(coord[,1], coord[,2], coord[,3], radius=0.03, color=coord[,4], alpha=1,
+ shininess=20); aspect3d(1, 1, 1)
> axes3d(col='black'); title3d("", "", "", "", "", col='black'); bg3d("white")
> # To save a snapshot of the graph, one can use the command rgl.snapshot("test.png").
```

19.5 Clustering with Other Algorithms

ChemmineR allows the user to take advantage of the wide spectrum of clustering utilities available in R. An example on how to perform hierarchical clustering with the `hclust` function is given below.

Create atom pair distance matrix:

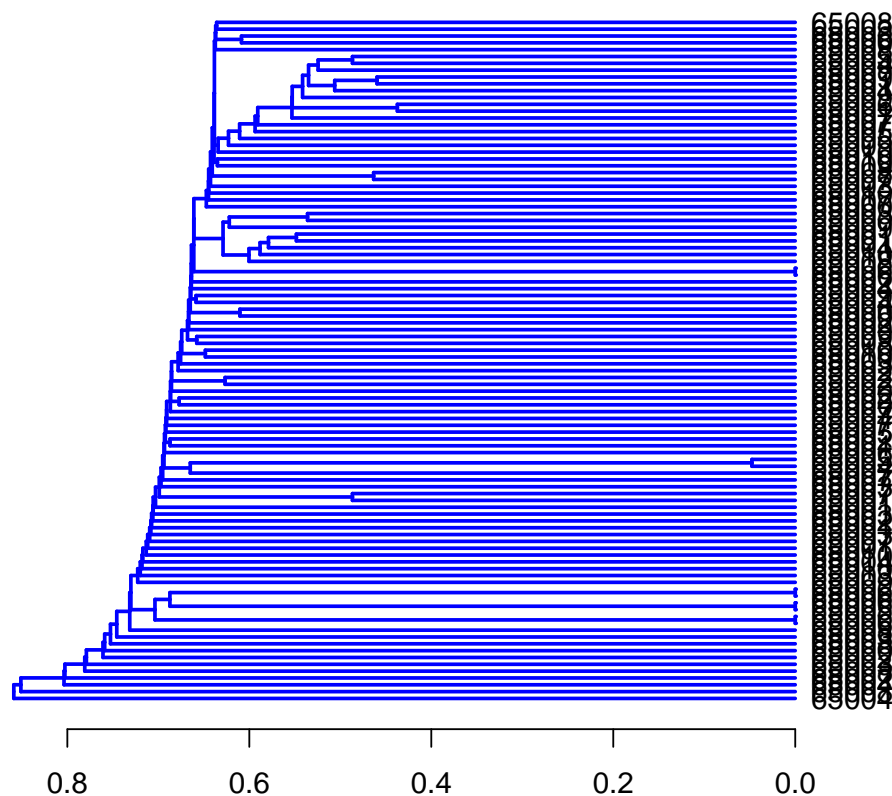
```
> dummy <- cmp.cluster(db=apset, cutoff=0, save.distances="distmat.rda", quiet=TRUE)
```

sorting result...

```
> load("distmat.rda")
```

Hierarchical clustering with `hclust`:

```
> hc <- hclust(as.dist(distmat), method="single")
> hc[["labels"]] <- cid(apset) # Assign correct item labels
> plot(as.dendrogram(hc), edgePar=list(col=4, lwd=2), horiz=T)
```

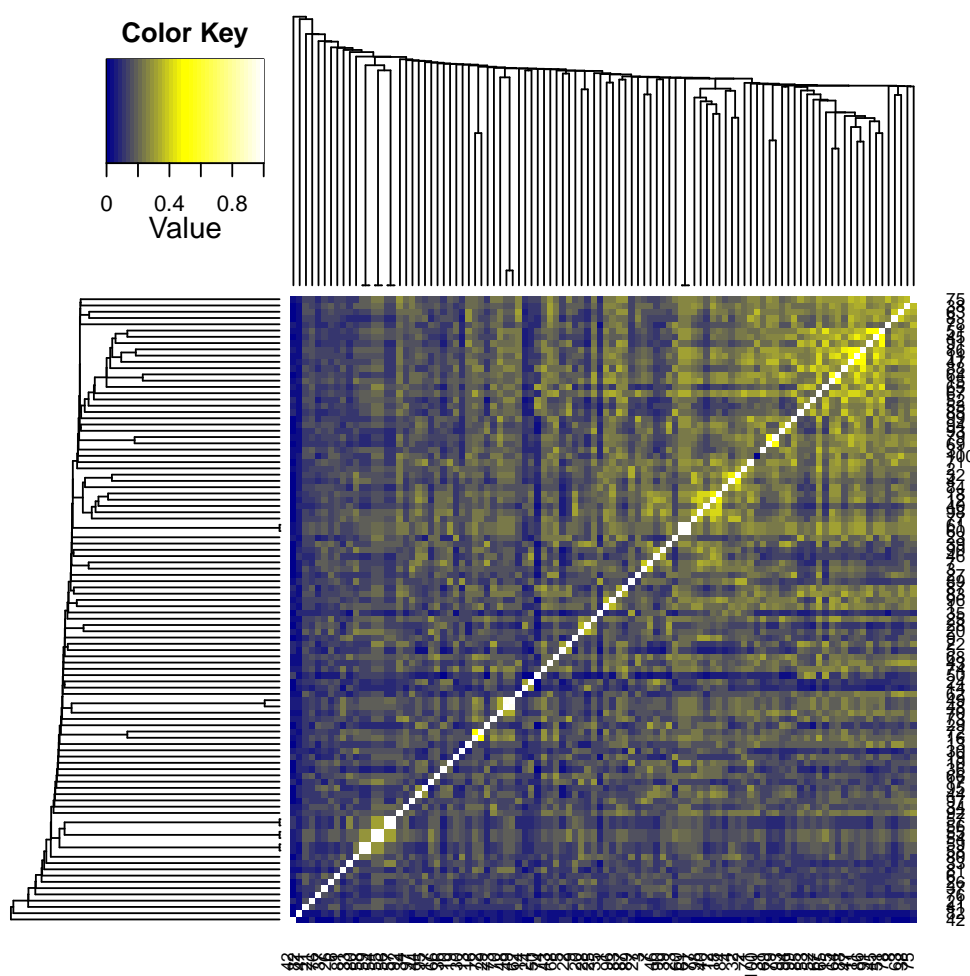


Instead of atom pairs one can use PubChem's fingerprints for clustering:

```
> simMA <- sapply(cid(fpset), function(x) fpSim(fpset[x], fpset, sorted=FALSE))
> hc <- hclust(as.dist(1-simMA), method="single")
> plot(as.dendrogram(hc), edgePar=list(col=4, lwd=2), horiz=TRUE)
```

Plot dendrogram with heatmap (here similarity matrix):

```
> library(gplots)
> heatmap.2(1-distmat, Rowv=as.dendrogram(hc), Colv=as.dendrogram(hc),
+           col=colorpanel(40, "darkblue", "yellow", "white"),
+           density.info="none", trace="none")
```



20 Searching PubChem

20.1 Get Compounds from PubChem by Id

The function `getIds` accepts one or more numeric PubChem compound ids and downloads the corresponding compounds from PubChem Power User Gateway (PUG) returning results in an *SDFset* container. The ChemMine Tools web service is used as an intermediate, to translate queries from plain HTTP POST to a PUG SOAP query.

Fetch 2 compounds from PubChem:

```
> compounds <- getIds(c(111,123))  
> compounds
```

20.2 Search a SMILES Query in PubChem

The function `searchString` accepts one SMILES string (Simplified Molecular Input Line Entry Specification) and performs a >0.95 similarity PubChem fingerprint search, returning the hits in an *SDFset* container. The ChemMine Tools web service is used as an intermediate, to translate queries from plain HTTP POST to a PubChem Power User Gateway (PUG) query.

Search a SMILES string on PubChem:

```
> compounds <- searchString("CC(=O)OC1=CC=CC=C1C(=O)O")
> compounds
```

20.3 Search an SDF Query in PubChem

The function `searchSim` performs a PubChem similarity search just like `searchString`, but accepts a query in an *SDFset* container. If the query contains more than one compound, only the first is searched.

Search an *SDFset* container on PubChem:

```
> data(sdfsamples); sdfset <- sdfsamples[1]
> compounds <- searchSim(sdfset)
> compounds
```

21 Version Information

```
> toLatex(sessionInfo())
```

- R version 3.0.3 (2014-03-06), i386-w64-mingw32
- Locale: LC_COLLATE=C, LC_CTYPE=English_United States.1252, LC_MONETARY=English_United States.1252, LC_NUMERIC=C, LC_TIME=English_United States.1252
- Base packages: base, datasets, grDevices, graphics, methods, stats, utils
- Other packages: ChemmineR 2.14.3, DBI 0.2-7, RSQLite 0.11.4, fmcsR 1.4.0, gplots 2.12.1, scatterplot3d 0.3-35
- Loaded via a namespace (and not attached): BiocStyle 1.0.0, KernSmooth 2.23-10, RCurl 1.95-4.1, bitops 1.0-6, caTools 1.16, digest 0.6.4, gdata 2.13.2, gtools 3.3.1, tools 3.0.3

22 Funding

This software was developed with funding from the National Science Foundation: [ABI-0957099](#), 2010-0520325 and IGERT-0504249.

23 References

- T W Backman, Y Cao, and T Girke. ChemMine tools: an online service for analyzing and clustering small molecules. *Nucleic Acids Res*, 39(Web Server issue):486–491, Jul 2011. doi: 10.1093/nar/gkr320. URL <http://www.hubmed.org/display.cgi?uids=21576229>.
- Y Cao, A Charisi, L C Cheng, T Jiang, and T Girke. ChemmineR: a compound mining framework for R. *Bioinformatics*, 24(15):1733–1734, Aug 2008. doi: 10.1093/bioinformatics/btn307. URL <http://www.hubmed.org/display.cgi?uids=18596077>.
- R.E. Carhart, D.H. Smith, and R. Venkataraghavan. Atom pairs as molecular features in structure-activity studies: definition and applications. *Journal of Chemical Information and Computer Sciences*, 25(2):64–73, 1985.
- X. Chen and C.H. Reynolds. Performance of Similarity Measures in 2D Fragment-Based Similarity Searching: Comparison of Structural Descriptors and Similarity Coefficients. *Journal of Chemical Information and Computer Sciences*, 42(6): 1407–1414, 2002.
- Th. Hanser, Ph. Jauffret, and G. Kaufmann. A new algorithm for exhaustive ring perception in a molecular graph. *Journal of Chemical Information and Computer Sciences*, 36(6):1146–1152, 1996. doi: 10.1021/ci960322f. URL <http://pubs.acs.org/doi/abs/10.1021/ci960322f>.

- R.A. Jarvis and E.A. Patrick. Clustering using a similarity measure based on shared near neighbors. *IEEE Transactions on Computers*, 22(11):1025–1034, 1973. URL http://davide.eynard.it/teaching/2012_PAMI/JP.pdf.
- N M O'Boyle, C Morley, and G R Hutchison. Pybel: a Python wrapper for the OpenBabel cheminformatics toolkit. *Chem Cent J*, 2(1):5–5, Mar 2008. doi: 10.1186/1752-153X-2-5. URL <http://www.hubmed.org/display.cgi?uids=18328109>.
- N M O'Boyle, M Banck, C A James, C Morley, T Vandermeersch, and G R Hutchison. Open Babel: An open chemical toolbox. *J Cheminform*, 3:33–33, 2011. doi: 10.1186/1758-2946-3-33. URL <http://www.hubmed.org/display.cgi?uids=21982300>.
- Y Wang, T W Backman, K Horan, and T Girke. fmcsR: Mismatch Tolerant Maximum Common Substructure Searching in R. *Bioinformatics*, Aug 2013. doi: 10.1093/bioinformatics/btt475. URL <http://www.hubmed.org/display.cgi?uids=23962615>.