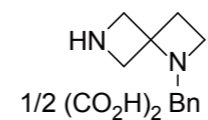


## Spirocyclic Compounds

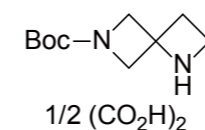
1351261



1223573-42-9

1-Benzyl-1,6-diazaspiro[3.3]heptane  
hemioxalate, 95%

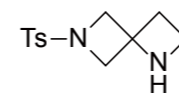
1351260



1431868-60-8

6-Boc-1,6-diazaspiro[3.3]heptane  
hemioxalate, 95%

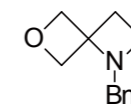
1590282



1223573-45-2

6-[(4-Methylphenyl)sulfonyl]-1,6-  
diazaspiro[3.3]heptane, 95%

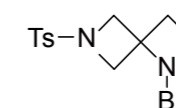
1556975



1223573-38-3

1-Benzyl-6-oxa-1-azaspiro  
[3.3]heptane, 95%

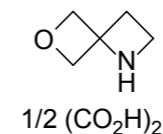
1565863



1223573-36-1

6-[(4-Methylphenyl)sulfonyl]-1-benzyl-  
1,6-diazaspiro[3.3]heptane, 95%

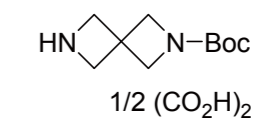
1238560



1046153-00-7

6-Oxa-1-azaspiro[3.3]heptane  
hemioxalate, 95%

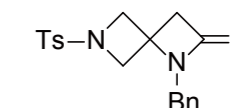
1238563



1041026-70-3

2-Boc-2,6-diazaspiro[3.3]heptane  
hemioxalate, 95%

1565862



1263296-91-8

6-[(4-Methylphenyl)sulfonyl]-1-benzyl-  
1,6-diazaspiro[3.3]heptan-2-one, 95%

# Spirocyclic Compounds

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
F: +852 2810 5033

E: [info@jk-scientific.com](mailto:info@jk-scientific.com)

# Spirocyclic Compounds

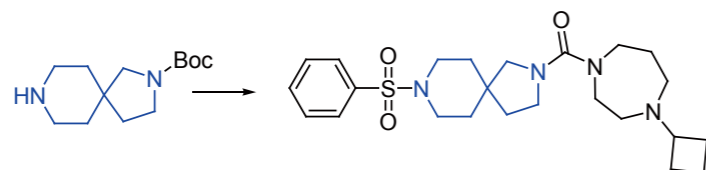
A spiro compound is a bicyclic organic compound with rings connected through just one atom. The rings can be different in nature or identical. The connecting atom, called the spiroatom, is most often a quaternary carbon ("spiro carbon").

The structural rigidity and feasibility for further structural elaboration through three-dimensional drug space have allowed spiro-bicyclic compounds to be applied in many bioactive molecules; this is one of the key strategies in modern medicinal chemistry.

J&K Scientific categorizes these novel spiro compounds based simply on their ring systems. For instance, 2-Oxa-6-azaspiro[3.4]octane () is one of the chemicals of the 5+4 system. We can provide all of the common kinds of spiro compound systems as outlined below:

## ■ [6+5]-Spirocyclic compounds:

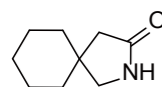
2-Boc-2,8-Diaza-spiro[4.5]decane has been used as the key intermediate in the synthesis of spirofused piperazine and diazepane amides. These compounds should provide useful starting points and tools to investigate further the importance of selective H<sub>3</sub> antagonists for drug therapy in a variety of potential disease states.



## References

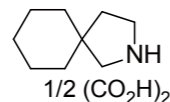
Brown, D. G.; Bernstein, P. R.; et al. J. Med. Chem., **2014**, 57, 733–758

511395



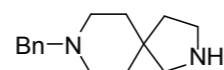
64744-50-9  
3-Azaspiro[4.5]decan-2-one, 98%

1175510



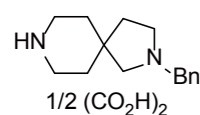
176-66-9  
2-Azaspiro[4.5]decane hemioxalate, 95%

912509



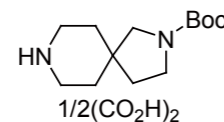
336191-15-2  
8-Benzyl-2,8-diazaspiro[4.5]decane, 95%

1549294



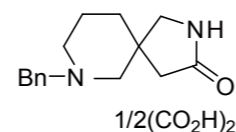
867009-61-8  
2-Benzyl-2,8-diazaspiro[4.5]decane hemioxalate, 95%

1711618



1086395-18-7  
7-Benzyl-2,7-diazaspiro[4.5]decane hemioxalate, 95%

1711617



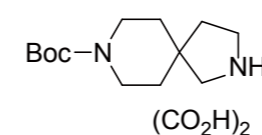
1312760-55-6  
7-Benzyl-2,7-diazaspiro[4.5]decan-3-one hemioxalate, 95%

1558751



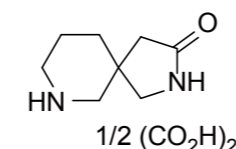
154495-69-9  
8-Benzyl-2,8-diazaspiro[4.5]decan-3-one, 95%

1139418



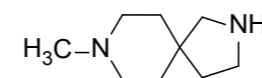
236406-39-6  
8-Boc-2,8-diazaspiro[4.5]decane oxalate, 95%

1553395



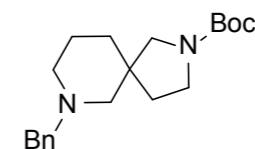
1158750-89-0  
2,7-Diazaspiro[4.5]decan-3-one hemioxalate, 95%

1117680



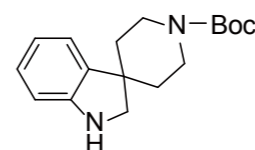
1158750-98-1  
8-Methyl-2,8-diazaspiro[4.5]decane, 95%

1549291



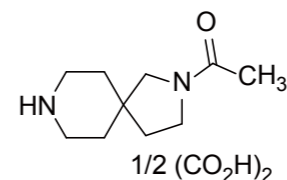
1245649-93-7  
2-Boc-7-benzyl-2,7-diazaspiro[4.5]decane, 95%

1019223



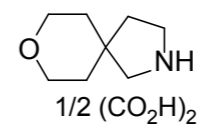
180465-84-3  
1'-Boc-spiro[indoline-3,4'-piperidine], 95%

1586814



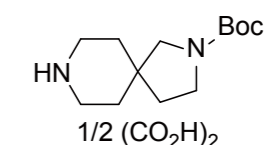
870082-43-2  
1-(2,8-Diazaspiro[4.5]decan-2-yl)ethanone hemioxalate, 95%

908679



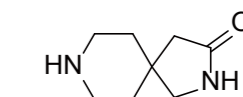
310-93-0  
8-Oxa-2-azaspiro[4.5]decane hemioxalate, 95%

328108



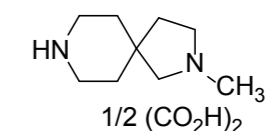
336191-17-4  
2-Boc-2,8-diazaspiro[4.5]decane hemioxalate, 95%

1117933



561314-57-6  
2,8-Diazaspiro[4.5]decan-3-one, 95%

1139419

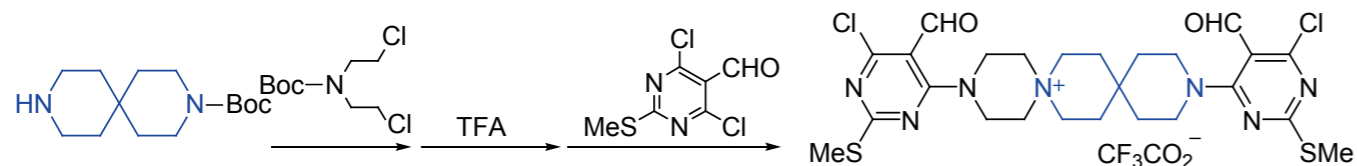


1061873-16-2  
2-Methyl-2,8-diazaspiro[4.5]decane hemioxalate, 95%

# Spirocyclic Compounds

## ■ [6+6]-Spirocyclic compounds:

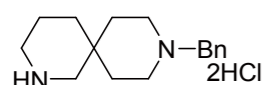
3,9-Diazaspiro[5.5]undecane has been used in the synthesis of the analogues of adhesamine. A study combined chemical, physicochemical, and cell biological experiments, using adhesamine and its analogues, to examine the mechanism by which this dumbbell-shaped, nonpeptidic molecule induces physiologically relevant cell adhesion. The results suggest that multiple adhesamine molecules cooperatively bind to heparan sulfate and induce its assembly, promoting clustering of heparan sulfate-bound syndecan-4 on the cell surface.



### References

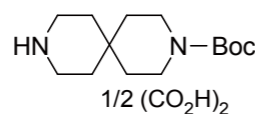
Takemoto, N.; Suehara, T.; et al. *J. Am. Chem. Soc.*, **2013**, 135, 11032–11039.

1574407



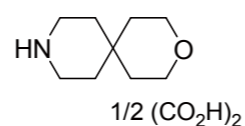
1198286-24-6  
9-Benzyl-2,9-diazaspiro[5.5]undecane dihydrochloride, 95%

536697



173405-78-2  
3-Boc-3,9-diazaspiro[5.5]undecane hemioxalate, 95%

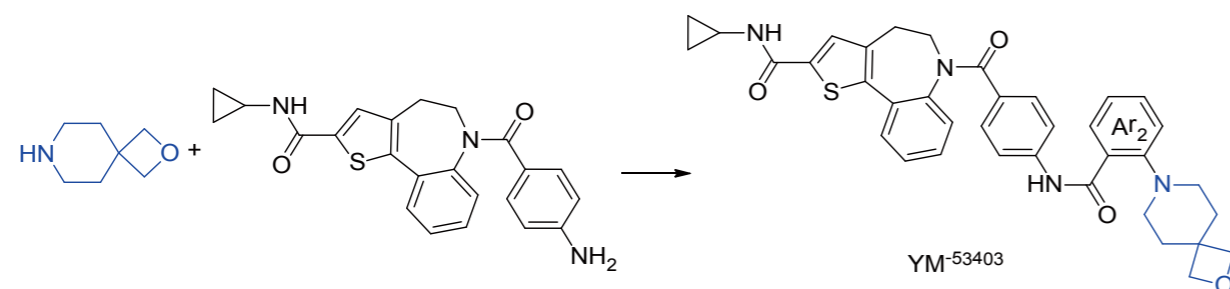
1139449



311-21-7  
3-Oxa-9-azaspiro[5.5]undecane hemioxalate, 95%

## ■ [6+4]-Spirocyclic compounds:

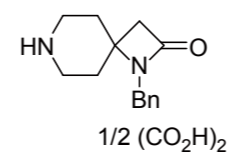
Modified by an oxetane spiro-fused piperidine, the derivatives of YM-53403 have been found to give significantly improved antiviral activity against RSV A2 and RSV B-WST; this discovery provides new knowledge that may pave the way towards effective RSV therapeutics and new tool compounds to interrogate RSV L protein function.



### References

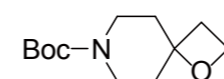
Xiong, H.; Foulk, M.; et al. *Bioorg. Med. Chem. Lett.*, **2013**, 23, 6789 – 6793.

1711613



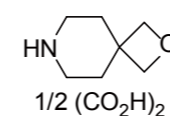
1415396-42-7  
1-Benzyl-1,7-diazaspiro[3.5]nonane hemioxalate, 95%

1494423



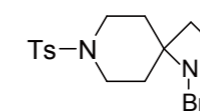
864684-96-8  
7-Boc-1-oxa-7-azaspiro[3.5]nonane, 96%

1548995



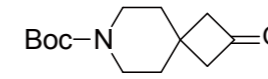
1379811-94-5  
2-Oxa-7-azaspiro[3.5]nonane hemioxalate, 95%

1711615



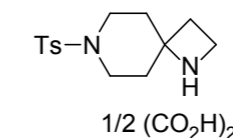
1-Benzyl-7-tosyl-1,7-diazaspiro[3.5]nonane, 95%

1350800



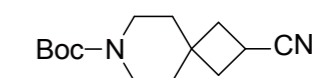
203661-69-2  
7-Boc-2-oxo-7-azaspiro[3.5]nonane, 95%

1711616



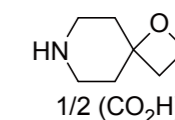
7-Tosyl-1,7-diazaspiro[3.5]nonane hemioxalate, 95%

1351181



203662-66-2  
7-Boc-2-cyano-7-azaspiro[3.5]nonane, 95%

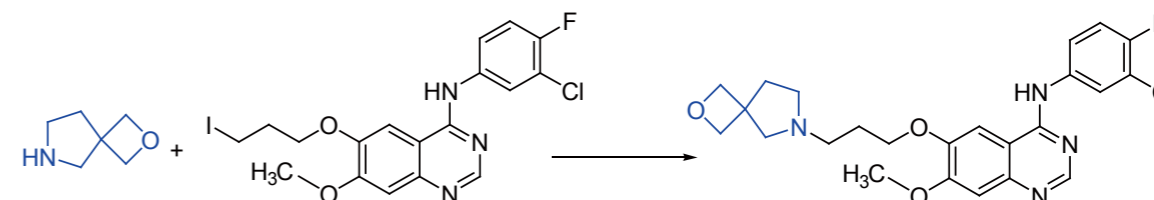
1581756



1408076-14-1  
1-Oxa-7-azaspiro[3.5]nonane hemioxalate, 97%

## ■ [5+4]-Spirocyclic compounds:

When 2-oxa-6-azaspiro[3.4]octane was substituted in 4-anilinoquinazoline derivative, higher EGFR inhibitory activities against two lung cancer cell lines (HCC827 and A549) were evaluated.



# Spirocyclic Compounds

## References

Zhao, F.; Lin, Z. H.; Wang, F.; et al. *Bioorg. Med. Chem. Lett.*, **2013**, 23, 5385-5388

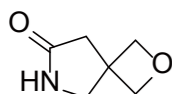
1581815



1392211-22-1

2-Benzyl-2,6-diazaspiro[3.4]octan-7-one, 95%

1238567

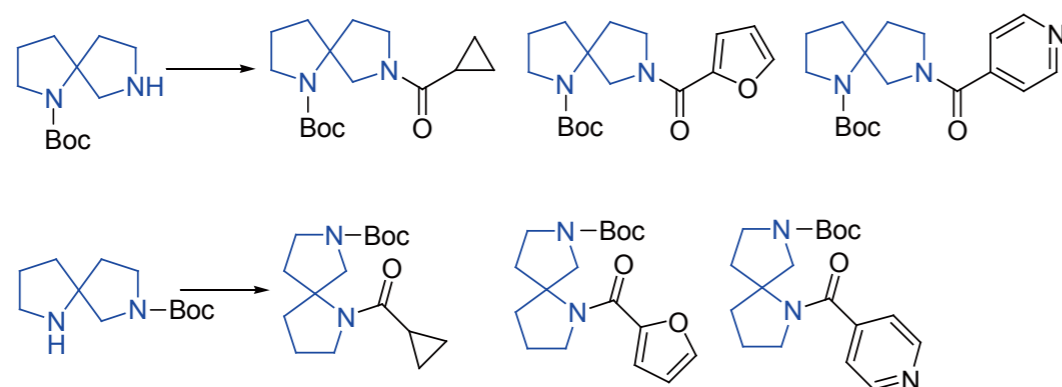


1207174-87-5

2-Oxa-6-azaspiro[3.4]octan-7-one, 95%

## ■ [5+5]-Spirocyclic compounds

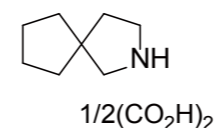
These spiro-bicyclic compounds have been used in the synthesis of the analogues of  $\alpha 4\beta 2$  nicotinic acetylcholine receptor agonist; They have been applied in the study of a potential treatment for cognitive deficits associated with psychiatric and neurological conditions.



## References

Mazurov, A. A.; Miao, L.; Bhatti, B. S.; et al. *J. Med. Chem.*, **2012**, 55, 9181- 9194

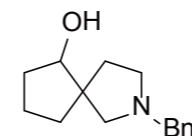
434240



1523617-88-0

2-Azaspiro[4.4]nonane hemioxalate, 95%

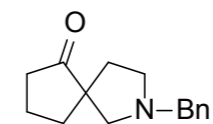
1762766



186202-97-1

6-Hydroxy-2-benzyl-2-azaspiro[4.4]nonane, 95%

1742307



160746-93-0

2-Benzyl-2-azaspiro[4.4]nonan-9-one, 95%

1586812

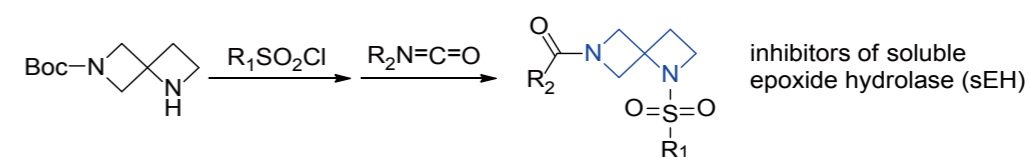


128244-01-9

1-Benzyl-1,7-diazaspiro[4.4]nonane hemioxalate, 95%

## ■ [4+4]-Spirocyclic compounds:

Some azetidine derivatives have been synthesized and used in therapy and/or prophylaxis, in particular to inhibitors of soluble epoxide hydrolase (sEH). The compounds are useful for treating disease states mediated by sEH, including genitourinary disease states, pain disease states, respiratory disease states, neurological disease states, immunological disease states, inflammatory disease states, cancer, nephropathy, stroke, endothelial dysfunction, prevention of ischemic events and end organ protection.



## References

Ceccarelli, S. M.; Guerot, c.; Knust, H. US 20130109668.