

Review

Selective Halogenation and Pseudohalogenation*

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Recent studies on addition of halogens, interhalogens, and pseudohalogens to alkenes and alkynes, substitution of hydrogen by halogens in alkane and benzene derivatives, and replacement of halogens of alkyl halides by thiocyanato, all of which proceed with high stereo- and/or regio-selectivities, are reviewed.

KEY WORDS: Halogenation/ Pseudohalogenation/ Alkenes/ Alkynes/
Arenes/

Some excellent reviews¹⁾ are available concerning the direct introduction of halogen or pseudohalogen species into organic substrates by addition or substitution. The general and theoretical aspects on these reactions can be found in most of these reviews. In this review article, therefore, some recent studies related to stereo- and/or regio-selective halogenation and pseudohalogenation are summarized concisely from synthetic viewpoint.

I. Addition of Halogens to Alkenes and Alkynes

It has been well-known that the addition of I_2 or Br_2 to aliphatic alkenes and alkynes in solvents of low polarity generally affords the *anti* adducts almost exclusively, whereas in the addition of Cl_2 involving a weakly bridged ion as the product-determining intermediate the adducts are formed with somewhat lower stereospecificity. For aryl-substituted alkenes, Br_2 adds in *anti* mode with considerable stereospecificity, but the addition of Cl_2 shows an excess of *syn* addition suggesting a carbenium-chloride ion pair intermediate.^{2,3)} Though an open vinyl cation intermediate is generally assumed for the halogenation of aryl-substituted alkynes, the reaction proceeds with considerable *anti* stereoselectivity except for the addition of Cl_2 , so far as the reaction is carried out in solvents of low polarity.^{5,6,7)} Recent stereochemical data on the halogenation of some alkenes and alkynes are shown in Table I. In the bromide-catalyzed bromination of olefins and acetylenes which probably proceeds through a termolecular mechanism, a considerable increase in the proportion of the *anti* adducts has been known. Recently the use of Br_2 -crown ether or Br_2 -pyridine complexes in the place of Br_2 is recommended to improve

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Table I. Stereochemistry of Halogenation of Some Alkenes and Alkynes with Halogens.

Substrate	Halogen	Solv. and Temp. (°C)	<i>anti</i> -Adduct (%) in Dihalide	Ref.
<i>t</i> -PhCH=CHMe	Br ₂	CCl ₄ , 2-5	88	2
<i>c</i> -PhCH=CHMe	Br ₂	CCl ₄ , 2-5	83	2
<i>t</i> -PhCH=CHMe	Cl ₂	CCl ₄ , 0-5	45 ^{a)}	3
<i>c</i> -PhCH=CHMe	Cl ₂	CCl ₄ , 0-5	32 ^{b)}	3
<i>t</i> -PhCH=CHMe	F ₂	CCl ₃ F, -78	31	4
<i>c</i> -PhCH=CHMe	F ₂	CCl ₃ F, -78	22	4
<i>n</i> -C ₆ H ₁₃ C≡CH	Br ₂	CHCl ₃ , 48	96	5
PrC≡CPr	Br ₂	CHCl ₃ , 20	100	5
EtC≡CEt	Cl ₂	CCl ₄ , 35	96	6
PhC≡CH	I ₂	CHCl ₃ , 60	100	7
PhC≡CPh	I ₂ +MeCO ₃ H	AcOH-Et ₂ O, 25	100	8
PhC≡CH	Br ₂	CHCl ₃ , 20	85	5
PhC≡CEt	Br ₂	CHCl ₃ , 20	86	5
PhC≡CH	Cl ₂	CCl ₄ , 0	49	6
PhC≡CEt	Cl ₂	CCl ₄ , 0	50	6

a) Other product, *c*-PhCH=C(Cl)Me. Dichloride: monochloride=84:16.b) Other product, *t*-PhCH=C(Cl)Me. Dichloride: monochloride=95:5.Table II. Effects of Solvents and Additives on Stereochemistry of Bromination of β -Methylstyrene and Phenylacetylene.

Substrate	Halogenating Agent	Solvent (ϵ)	<i>anti</i> -Adduct (%) in Dibromide	Ref.
<i>t</i> -PhCH=CHMe	Br ₂	C ₆ H ₁₂ (2.0)	92	9
	Br ₂ -DBC ^{a)}	C ₆ H ₁₂	100	9
	Br ₂	AcOH (6.2)	83 ^{b,c)}	10
	Br ₂ -LiBr	AcOH	95 ^{b,d)}	10
	Br ₂	CH ₂ Cl ₂ (8.9)	89	11
	Br ₂ -pyridine	CH ₂ Cl ₂	98	11
	Br ₂ -DBC	CH ₂ Cl ₂	100	9
	Br ₂	C ₆ H ₁₂	85	9
<i>c</i> -PhCH=CHMe	Br ₂ -DBC	C ₆ H ₁₂	100	9
	Br ₂	AcOH	73 ^{b,c)}	10
	Br ₂ -LiBr	AcOH	95 ^{b,d)}	10
	Br ₂	CH ₂ Cl ₂	75	11
	Br ₂ -PyHBr	CH ₂ Cl ₂	100	11
	Br ₂ -LiClO ₄	AcOH	52 ^{e)}	12
PhC≡CH	Br ₂ -LiBr	AcOH	>99	12

a) Dibenzo-18-crown-6.

b) Other product, PhCH(OAc)CH₂Br.c) Dibromide: bromoacetate=*ca.* 75-80: 20-25.d) Dibromide: bromoacetate=*ca.* 85: 15.e) Other products; *E*- & *Z*-Ph(OAc)C=CHBr and PhCOCHBr₂.
Dibromide: other products=73: 27.

the *anti* selectivity in the bromination of aryl-substituted alkenes and alkynes (Table II).^{9,11} Further, it is found that the radical chlorination of alkyl and aryl alkynes with Cl_2 , SO_2Cl_2 , or benzene iododichloride appears to be efficient for the selective formation of the corresponding *E*-dichlorides (Table III).^{6,13,14} It should be noted here that the possibility of a change in the ratio of the *E*- and *Z*-isomer due to prolonged reaction is sometimes present.

 Table III. Stereochemistry of Radical Chlorination of β -Methylstyrene and Some Alkynes.

Substrate	Halogenating Agent	Solv. and Temp. ($^{\circ}\text{C}$)	<i>anti</i> -Adduct (%)		Ref.
			in Product	in Equilibrium	
<i>t</i> -PhCH=CHMe	PhICl ₂ ^a	CHCl ₃ , 60	94	—	13
<i>c</i> -PhCH=CHMe	PhICl ₂ ^a	CHCl ₃ , 60	6	—	13
<i>n</i> -C ₆ H ₁₃ C $\equiv$ CH	Cl ₂ , <i>h</i> ν	CCl ₄ , 0	94		6
	PhICl ₂ ^a	CHCl ₃ , 60	94	53	6
	SO ₂ Cl ₂	C ₆ H ₆ , 80	93		14
PhC $\equiv$ CH	Cl ₂ , <i>h</i> ν	CCl ₄ , 0	83		6
	PhICl ₂ ^a	CHCl ₃ , 60	89	14	6
	SO ₂ Cl ₂	C ₆ H ₆ , 80	85		14
PhC $\equiv$ CEt	Cl ₂ , <i>h</i> ν	CCl ₄ , 0	75		6
	SO ₂ Cl ₂	C ₆ H ₆ , 80	75		14
PhC $\equiv$ CPh	PhICl ₂ ^a	CHCl ₃ , 60	76	28	6

a) Azobisisobutyronitrile (radical initiator) is added.

Compared with halogenation by molecular halogens, some characteristics have been found in halogen addition by metal halides. For example, SbCl_5 or MoCl_5 reacts with alkenes and alkynes to afford the *syn* adducts predominantly.^{15,16,21} On the contrary, the favorable formation of the *E*-dihalides from alkynes is found in the case of CuBr_2 or CuCl_2 .^{5,20} Recently, an efficient but rather complex *in situ* Mo(VI) reagent for the *syn* chlorination of alkenes has been developed.¹⁷ Some data on the halogenation of alkenes and alkynes using metal halides are tabulated in Table IV. Halogenation by metal halides appears to be synthetically useful because highly stereoselective dichlorides or dibromides can be easily prepared.

A comparison of the halogenation of butadiene with Br_2 and Cl_2 has been carried out. The effect of solvent polarity on regioselectivity is somewhat significant in the bromination, whereas it is insensitive in the chlorination.²² Butadiene reacts with Br_2 to give the 1,4-adduct with *trans* configuration mainly, but when Br_2 -pyridine complex is used as a brominating agent the 1,2-adduct is formed predominantly.¹¹ On the other hand, its chlorination by Cl_2 yields a mixture of almost equal amounts of the 1,2- and 1,4-adducts, while the reaction with SbCl_5 at low temperatures is highly stereoselective toward the formation of the thermodynamically least stable product, *cis*-1,4-dichloro-2-butene.²³ Several data are shown in Table V. A lower regioselectivity in the Cl_2 addition compared to the Br_2 addition may be due to an extensive dispersal of positive charge into the vinylic system.²² The

Table IV. Stereochemistry of Halogenation of Some Alkenes and Alkynes with Metal Halides.

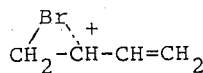
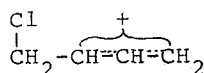
Substrate	Metal Halide	Solv. and Temp. (°C)	Yield (%)	anti-Adduct (%) in Product	Ref.
<i>t</i> -MeCH=CHMe	SbCl ₅	CCl ₄ , 76	96	18	15
	MoCl ₅	CCl ₄ , 76	85	16	16
<i>c</i> -MeCH=CHMe	SbCl ₅	CCl ₄ , 76	98	16	15
	MoCl ₅	CCl ₄ , 76	92	15	16
<i>t</i> -PrCH=CHPr	(Bu ₄ N) ₄ Mo ₈ O ₂₈ + AcCl	CH ₂ Cl ₂ , 25	91	2	17
<i>cyclo</i> -C ₆ H ₁₀	MoCl ₅	CCl ₄ , -10~0	70	2	18
<i>t</i> -PhCH=CHMe	SbCl ₅	CCl ₄ , -10~0	63	25	19
<i>c</i> -PhCH=CHMe	SbCl ₅	CCl ₄ , -10~0	67	0	19
<i>n</i> -C ₆ H ₁₃ C≡CH	CuBr ₂	MeCN, 25	73	100	5
PrC≡CPr	CuBr ₂	MeCN, 25	65	100	5
PhC≡CH	CuBr ₂	MeCN, 25	77	97	5
PhC≡C- <i>i</i> -Pr	CuBr ₂	MeCN, 25	60	91	5
PhC≡C- <i>t</i> -Bu	CuBr ₂	MeCN, 25	100	9	5
<i>n</i> -C ₆ H ₁₃ C≡CH	CuCl ₂ -LiCl	MeCN, 82	83	92	20
PrC≡CPr	CuCl ₂ -LiCl	MeCN, 82	42	98	20
	MoCl ₅	CH ₂ Cl ₂ , 25	37	3	18
PhC≡CH	CuCl ₂ -LiCl	MeCN, 82	71	94	20
PhC≡CEt	SbCl ₅	CCl ₄ , 25	32	16	21
PhC≡C- <i>t</i> -Bu	CuCl ₂ -LiCl	MeCN, 82	94	21	20
	SbCl ₅	CCl ₄ , 76	55	0	21

Table V. Product Distribution of Halogenation of Butadiene.

Halogenating Agent	Solv. and Temp. (°C)	Product Distribution (%)			1, 2-Add. 1, 4-Add.	Ref.
		1,2-	<i>t</i> -1, 4-	<i>c</i> -1, 2-		
Br ₂	<i>n</i> -C ₅ H ₁₂ , -10	69	31	—	2.2	22
Br ₂	CCl ₄ , -10	57	43	—	1.3	22
Br ₂	CH ₂ Cl ₂ , -10	28	72	—	0.35	22
Br ₂ -Py	CH ₂ Cl ₂ , 0~5	85	15	—	5.7	11
Br ₂ -Py+Py	CH ₂ Cl ₂ , 0~5	96	4	—	24	11
Cl ₂	<i>n</i> -C ₅ H ₁₂ , -10	55	45	— ^{a)}	1.2	22
Cl ₂	CCl ₄ , -10	57	43	—	1.3	22
Cl ₂	CH ₂ Cl ₂ , -20~-10	54	45	<1	1.2	23
SbCl ₅	CH ₂ Cl ₂ , -26~-13	35	25	40	0.54	23
SbCl ₅	CH ₂ Cl ₂ , -22~-12 ^{b)}	25	51	24	0.33	23
CuCl ₂	CH ₃ CN, 60	15	80	5	0.18	33

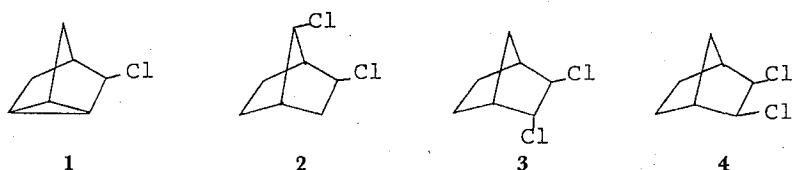
a) The isomer ratio of dichlorobutenes in equilibrium (neat, at 60°C), 1, 2-: *t*-1, 4-: *c*-1, 4- = 17: 77: 6 (Ref. 23).

b) Reverse addition mode. Diene was added to SbCl₅ solution.

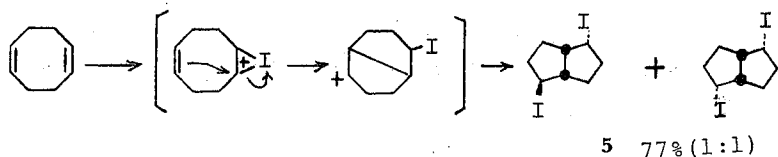


addition of Br_2 or Cl_2 to cyclopentadiene, a conjugated cyclic diene, affords the *syn*-1,4-adduct favorably,²⁴⁾ but bromination by Br_2 -pyridine complex yields the *anti*-1,2-adduct almost exclusively.¹¹⁾

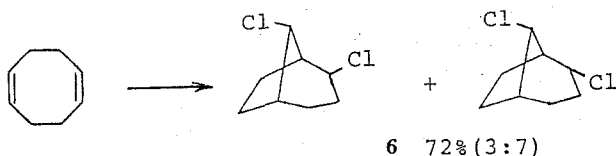
It has been known already that the main products in the ionic chlorination of norbornene with Cl_2 are nortricyclyl chloride (**1**) and *exo*-2-*syn*-7-dichloronorbornane (**2**), and no appreciable formation of the 2,3-dichloro isomer is observed. A recent study shows that treatment with a mixture of CuCl_2 and LiCl in MeCN or with MoCl_5 or VCl_5 in CCl_4 affords mainly *trans*- or *cis*-2,3-dichloronorbornane (**3**, **4**), respectively, in good yields.²⁵⁾ Interestingly, radical chlorination by SO_2Cl_2 or benzene iododichloride in CCl_4 also gives the *trans*-2,3-isomer favorably.^{26,27)}



The halogenation of cyclic monoolefins and unconjugated cyclic dienes with molecular halogens is known to proceed in usual mode, *i.e.*, 1,2-*anti*-addition. However, it is found that in the chlorination of *cis,cis*-1,5-cyclooctadiene with MoCl_5 in CCl_4 , or with SO_2Cl_2 or PCl_5 in refluxing CCl_4 the *cis*-1,2-dichloride is formed predominantly.^{26,28)} Further, though the reaction with Br_2 or Cl_2 gives the corresponding *trans*-1,2-dihalides, a similar treatment with I_2 in CCl_4 results in the formation of isomeric 2,6-diiodobicyclo[3.3.0]octanes (**5**) in good yields.²⁹⁾ A similar



transannular cyclization is known in the iodination of *cis*, *trans*-cyclodecadiene.³⁰⁾ In the chlorination of the above diene and *cis*-cyclooctene with SbCl_5 in CCl_4 at low temperature, the formation of unusual products is reported.³¹⁾ A slow addition of SbCl_5 to the diene gives mainly a mixture of *exo*- and *endo*-2-*anti*-8-dichlorobicyclo[3.2.1]octanes (**6**), which is probably formed by a transannular cyclization and its subsequent isomerization. Here, the reverse addition affords usual transannular



products (**5**, $\text{I} \rightarrow \text{Cl}$). In the case of *cis*-cyclooctene *cis*-1,4-dichlorocyclooctane is formed predominantly and this can be explained by a transannular 1,5-hydride

shift.

Some recent results relating to the stereo- and/or regio-chemistry of mixed halogenation of olefins and acetylenes are summarized in Table VI. Various interhalogen compounds are known, but they are usually unstable except for ICl. Therefore the reaction is generally carried out under the *in situ* generation of interhalogens. Here it should be noted that the use of the parent halogen is sometimes accompanied by the formation of homohalogenated product. For example, in the preparation of *vic*-bromochlorocyclohexane from cyclohexene, a considerable increase in the product selectivity has been observed, when KBr-CuCl₂ combination is used as a halogenating agent instead of Br₂-CuCl₂ couple.³³⁾ In the iodochlorination of olefins, it is revealed that I₂-CuCl₂ or I₂-SbCl₅ system is very often superior to ICl in the yield and regioselectivity of the products, and that I₂-SbCl₅ couple is especially suitable for the halogenation of olefins bearing electron-withdrawing groups.³²⁾ Recently, a new iodofluorinating agent, MeIF₂, prepared from MeI and XeF₂ with

Table VI. Stereo- and Regio-chemistry of Mixed Halogenation of Some Alkenes and Alkynes.

Substrate	Halogenating Agent	Solv. and Temp. (°C)	Yield (%)	<i>anti</i> -Adduct in Product (%)	M ^{a)} in Product (%)	Ref.
BuCH=CH ₂	ICl	CCl ₄ , 30	75		66	32
	I ₂ -CuCl ₂	MeCN, 25	91		80	33
CH ₂ =CHCl	I ₂ -CuCl ₂	MeCN, 50	81		73	33
CH ₂ =CHOAc	I ₂ -CuCl ₂	MeCN, 25	83		100	33
CH ₂ =CHCO ₂ Et	I ₂ -SbCl ₅	CCl ₄ , 76	86		18	32
<i>t</i> -EtOOCCH=CHCO ₂ Et	I ₂ -SbCl ₅	CCl ₄ , 76	60	100	—	32
PhCH=CH ₂	KI-CuCl ₂	MeCN, 80	75		100	33
<i>n</i> -C ₆ H ₁₃ CH=CH ₂	I ₂ -F ₂	CFCl ₃ , -75	70		100	34
BuCH=CH ₂	BrCl	CCl ₄ , 25	96		61	35
CH ₂ =CHOAc	Br ₂ -SbCl ₅	CCl ₄ , 5	79		100	36
CH ₂ =CHCO ₂ Me	Br ₂ -Cl ₂	CH ₂ Cl ₂ , 25	89		17	37
<i>t</i> -EtOOCCH=CHCO ₂ Et	Br ₂ -SbCl ₅	CCl ₄ , 5	66	100	—	36
PhCH=CH ₂	Br ₂ -SbCl ₅	CCl ₄ , 0	87		100	36
	Br ₂ -AgF	C ₆ H ₆ , 0~25	71		100	38
BuCH=CH ₂	NCS ^{b)} -HF	HF-Py-Sulfolane, 25	40		100	39
<i>n</i> -C ₆ H ₁₃ C≡CH	ICl	MeCN, 82	50	100	100	20
	I ₂ -CuCl ₂	MeCN, 82	84	100	100	20
PhC≡CH	ICl	MeCN, 82	52	95	100	20
	I ₂ -CuCl ₂	MeCN, 82	97	100	100	20
PhC≡CEt	I ₂ -CuCl ₂	MeCN, 82	100	100	100	20
PhC≡CMe	MeIF ₂	MeI-HF, 25	70	0	100	40
BuC≡CH	BrCl	CCl ₄ , 25	79	100	90	35
	NBA ^{c)} -HF	HF-Et ₂ O, -78~-20	48	95	100	41
EtC≡CEt	NBA-HF	HF-THF, -78~0	28	78	—	41

a) M; Markovnikov addition product. b) N-Chlorosuccinimide. c) N-Bromoacetamide.

HF catalyst, has been developed.⁴⁰⁾ This reacts with aryl-substituted alkenes to form the expected Markovnikov adducts with high *anti*-stereospecificity, while with aryl-substituted alkynes it affords only the *syn* adducts as the kinetically controlled products.⁴⁰⁾

II. Addition of Pseudohalogens to Alkenes and Alkynes

Compared to halogens, thiocyanogen is known to be a weak electrophile. Thus, under heterolytic conditions, it reacts with olefins in solvents of low polarity to form two *anti*-stereospecific products, *i.e.*, α , β -dithiocyanates and α -isothiocyanato- β -thiocyanates, in which the latter formation is favored,^{42,43)} while it does not react with acetylenes. In the reaction with olefins, relative yields of dithiocyanates can be increased by the addition of catalytic amounts of Fe or Fe(SCN)₃.⁴²⁾ On the contrary, under homolytic conditions, both alkenes and alkynes react with thiocyanogen readily. Simple alkenes afford the corresponding dithiocyanates with non-stereospecificity, while alkynes yield dithiocyanatoalkenes with high *anti*-stereoselectivity.^{44,45)} As in the case of dihalogenoalkenes, dithiocyanatoalkenes are also susceptible to subsequent thermodynamically-controlled isomerization. Several data are given in Table VII.

Table VII. Heterolytic and Homolytic Addition of Thiocyanogen to Some Alkenes and Alkynes.

Substrate	Solv. and Temp. (°C)	Yield (%)	Product Distribution (%)		Ref.
			S-S ^{a)}	S-N ^{a)}	
			(anti-Adduct %)		
<i>t</i> -EtCH=CH <i>Et</i>	MeCN, 25 ^{b)}	37	69(100)	31(100)	42
	C ₆ H ₆ , 25 ^{b)}	48	37(96)	63(98)	42
	C ₆ H ₆ , 25 ^{b,c)}	92	99.8(99)	0.2(100)	42
Me ₂ C=CMe ₂	C ₆ H ₆ , 25 ^{b)}	90	33(100)	67(100)	43
PhCH=CH ₂	C ₆ H ₆ , 25 ^{b)}	66	67	33	43
<i>t</i> -MeCH=CHMe	C ₆ H ₆ , <i>h</i> ν , 20~30	99	100(60)		44
<i>c</i> -MeCH=CHMe	C ₆ H ₆ , <i>h</i> ν , 20~30	94	100(39)		44
PhCH=CH ₂	C ₆ H ₆ , <i>h</i> ν , 20~30	99	100		44
HC $\equiv$ CH	C ₆ H ₆ , <i>h</i> ν , 20~25	90	100(95)		45
EtC $\equiv$ CEt	C ₆ H ₆ , <i>h</i> ν , 20~25	97	100(90)		45
PhC $\equiv$ CMe	C ₆ H ₆ , <i>h</i> ν , 20~25	90	100(89)		45
PhC $\equiv$ CPh	C ₆ H ₆ , <i>h</i> ν , 20~25	90	100(94)		45

a) S-S: Dithiocyanate. S-N: Isothiocyanatothiocyanate.

b) 2,6-Di-*t*-butyl-4-methylphenol (radical inhibitor) was added; in darkness.

c) Fe(SCN)₃ was added as catalyst.

It has been known that halogen azides can add to alkenes by either an ionic or free-radical mechanism. From the electronegativity trend $I < N_3 \approx Br < Cl$, it would be expected that the preference for an ionic mechanism would be $IN_3 > BrN_3 > ClN_3$. This is confirmed by comparison of the halogen azide addition to styrene under various conditions (Table VIII).⁴⁶⁾ Here, the Markovnikov adducts are formed by ionic route and the *anti*-Markovnikov adducts by radical pathway.

Table VIII. Heterolytic and Homolytic Addition of Halogen Azides to Styrene.

Reagent	Solv. and Temp. (°C)	Product Distribution (%)		Ref.
		M ^{a)}	aM ^{b)}	
IN ₃ (ICl+NaN ₃)	MeCN, 0	100	0	46
	<i>n</i> -C ₅ H ₁₂ , N ₂ , 0	61	39	46
BrN ₃ (Br ₂ +NaN ₃)	MeNO ₂ , N ₂ , 0	>96	<4	46
	<i>n</i> -C ₅ H ₁₂ , N ₂ , 0	0	100	46
ClN ₃ (Cl ₂ +NaN ₃)	MeNO ₂ , O ₂ , 0	48	17 ^{c)}	46
	<i>n</i> -C ₅ H ₁₂ , air, 0	0	100	46

a) Markovnikov adduct.

b) *anti*-Markovnikov adduct.

c) Other product, PhCH=CHCl (relative yield, 23%).

It is interesting that the nature of the reaction (*i.e.*, ionic or radical) is readily changed in the case of BrN₃.

In ionic media, ISC_N is polarized in the manner I^{δ+}-SC_N^{δ-}, while BrSC_N and ClSC_N are in the reverse direction, *i.e.*, X^{δ-}-SC_N^{δ+} (X=Br, Cl). When metal thiocyanates are used as a thiocyanogen source in the iodothiocyanation of olefins, product distribution varies significantly depending upon the metal salts. When KSCN is used *vic*-iodothiocyanoates by *anti* addition are formed predominantly, while the use of TISCN or Hg(SCN)₂ results in the favorable formation of *vic*-iodoisothiocyanoates.^{47,48)} The halogenoselenocyanation of olefins can be readily accomplished in MeCN by using a redox system of KSeCN and CuX₂ (X=Br or Cl) as an *in situ* XSeCN reagent. The reaction proceeds with Markovnikov-type regioselectivity and with a high *anti*-stereospecificity to form halogenoselenocyanates exclusively.⁵⁰⁾ Various data are shown in Table IX.

Table IX. Addition of Halogen Thiocyanates and Selenocyanates to Some Alkenes.

Substrate	Reagent	Solv. and Temp. (°C)	Yield (%)	Product Distribution (%)		Ref.
				X-YCN ^{a)}	X-NCY ^{a)}	
<i>cyclo</i> -C ₆ H ₁₀	I ₂ -KSCN	CHCl ₃ , 20	~100	96	4	47
	I ₂ -TISCN	CHCl ₃ -sulfolane, 0	75	20	80	47
	I ₂ -Hg(SCN) ₂	Et ₂ O, 20	80 ^{b)}	17	83	48
	I ₂ -(SCN) ₂	C ₆ H ₆ , 20	72 ^{b)}	26	74	48
<i>t</i> -EtCH=CHEt	I ₂ -KSCN	Et ₂ O, 20	63	95	5	48
	I ₂ -Hg(SCN) ₂	C ₆ H ₆ , 20	84	12	88	48
<i>cyclo</i> -C ₆ H ₁₀	Br ₂ -TISCN	CHCl ₃ , 25	51 ^{c)}	100	0	49
	CuBr ₂ -KSeCN	MeCN, 82	54	100	0	50
	CuCl ₂ -KSeCN	MeCN, 82	69	100	0	50
PhCH=CH ₂	CuBr ₂ -KSeCN	MeCN, 82	58	100	0	50
	CuCl ₂ -KSeCN	MeCN, 82	83	100	0	50

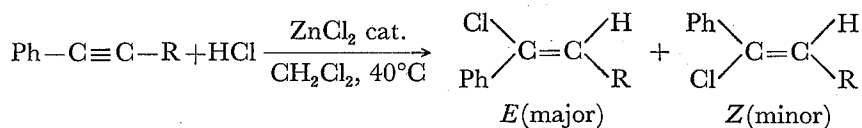
a) X-YCN: Halogenothiocyanoate (Y=S) or halogenoselenocyanate (Y=Se). X-NCY: Halogenoisothiocyanoate (Y=S) or halogenoisoselenocyanate (Y=Se). All products consist of *anti*-adducts entirely or mostly.

b) Other product, Dithiocyanate (1%).

c) Other products, Dibromide (7%) and dithiocyanate (15%).

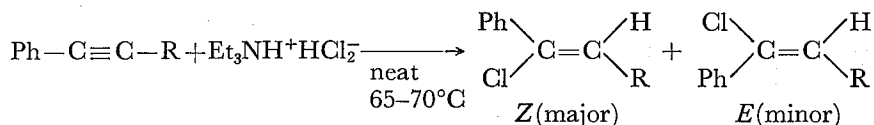
III. Introduction of Halogen or Pseudohalogen into Aliphatic Compounds by Other Reactions

The stereochemistry of HCl addition to phenylacetylenes has recently been studied. The addition of HCl in CH_2Cl_2 , in the presence of ZnCl_2 , proceeds with high *syn*-stereoselectivity, indicating the initial formation of an open vinyl cation intermediate and the subsequent preferential attack of Cl^- from the less hindered



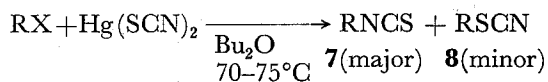
$\text{R}=\text{D}$ 85% ($E/Z=7/3$); $\text{R}=\text{Et}$ 40% ($E/Z=8/2$); $\text{R}=t\text{-Bu}$ 79% (only *E*)

site.⁵¹⁾ On the other hand, the hydrochlorination by $\text{Et}_3\text{NH}^+\text{HCl}_2^-$ without catalyst yields the *anti*-adducts predominantly, probably through a trimolecular mechanism.⁵²⁾



$\text{R}=\text{H}$ 73%; $\text{R}=\text{Me}$ 36% ($Z/E=89/11$)

The reaction of alkyl and aralkyl halides with KSCN in dipolar aprotic solvents is well-known as a preparative method of the corresponding thiocyanates. Based on the HASB principle, it would be expected that the predominant formation of isothiocyanate is achieved by a suitable choice of metal thiocyanates and/or solvents. In fact, it is found that when secondary or tertiary halides are treated with $\text{Hg}(\text{SCN})_2$ in solvents of low polarity (*e.g.*, *n*-hexane or di-*n*-butyl ether), the corresponding isothiocyanates are obtained with high selectivity.⁵³⁾ A mechanism



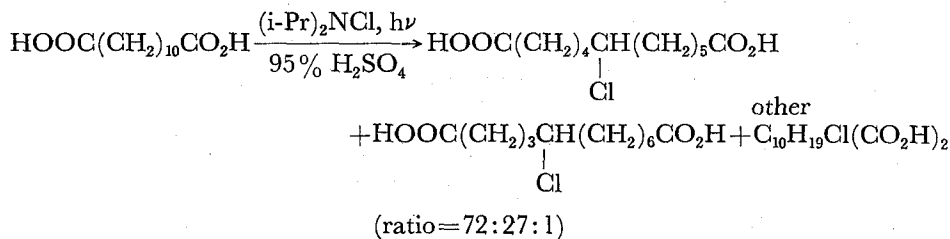
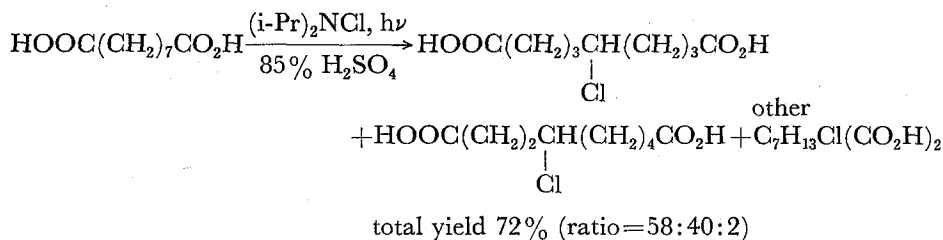
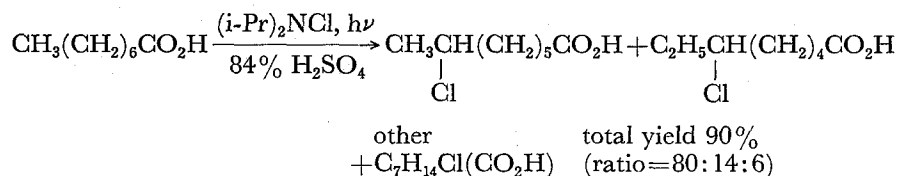
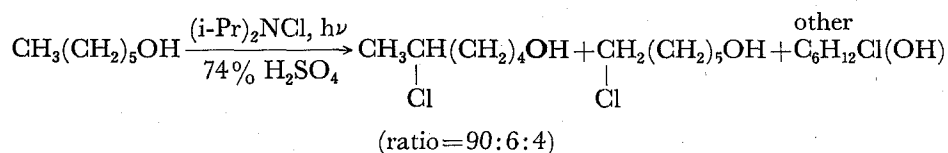
$\text{X}=\text{I}$, $\text{R}=i\text{-Pr}$ 72% ($7/8=85/15$); $\text{X}=\text{Br}$, $\text{R}=t\text{-Bu}$ 68% ($7/8=96/4$);

$\text{X}=\text{Br}$, $\text{R}=\text{PhCHMe}$ 87% ($7/8=99/1$)

involving the initial Hg-catalyzed ionization of alkyl halide and simultaneous formation of the complex ion $[\text{XHg}(\text{SCN})_2]^-$, followed by the attack of the terminal N site of this anion to alkyl cation, is proposed.

The introduction of halogen species at a definite position of long *n*-alkyl chain seems to be principally difficult, but the following finding is quite attractive from synthetic viewpoint. When $\text{C}_6\text{-C}_8$ aliphatic alcohols and acids are treated with sterically hindered *N*-chlorodiisopropylamine in conc. H_2SO_4 under irradiation, the chlorination occurs at the $\omega-1$ position of the substrates with high selectivity.⁵⁴⁾ An extension of this study to $\text{C}_8\text{-C}_{12}$ dicarboxylic acids has also been reported.⁵⁵⁾ In these reactions, an electrostatic repulsion between the protonated group (*e.g.*,

COOH_2^+) and the attacking radical cation (R_2NH^+) would be operative.



IV. Halogenation and Pseudohalogenation of Benzene Nucleus

Phenols, phenyl ethers, and anilines are readily halogenated, and usual halogenation by molecular halogens generally affords a mixture of *o*- and *p*-halogenated, and/or polyhalogenated compounds. A redox combination $\text{I}_2\text{-CuCl}_2$ has been developed as a simple and gentle iodinating reagent for aromatics, and in the iodination of phenol, anisole, and *N,N*-dimethylaniline with this reagent only the *para*-iodo-substituted products are formed.⁵⁶⁾ For the introduction of I into the *ortho*-position of phenols, $\text{I}_2\text{-TIOAc}$ is found to be appropriate.⁵⁷⁾ However, for anisole and anilines, this reagent leads to only *para*-substitution. Though the predominant *para*-bromination of phenols with Br_2 in various solvents has been known, it is reported that in CCl_4 or CHCl_3 solvent *N*-bromosuccinimide (NBS) brominates phenols in the *ortho*-position with high selectivity, but in polar aprotic solvents the reaction is *para*-selective.⁵⁸⁾ The use of NBS in DMF is quite useful for the selective monobromination of reactive aromatics including polynuclear hydrocarbons.⁵⁹⁾ Recently, it is also found that, in HF-SbF_5 , *p*-cresol and xylenols react with Br_2 to give the corresponding *meta*-bromo-substituted compounds in good yields, indicating the bromination of the O-protonated substrate.⁶⁰⁾ This method is not applicable to

Selective Halogenation and Pseudohalogenation

phenol, *o*- and *m*-cresol, all of which afford only the *para*-substitution products. In contrast to the halogenation of phenol with I₂ and Br₂, the effect of solvent on the product ratio appears to be significant in the chlorination by Cl₂. For example, the *o*:*p* ratio is 0.49 (in EtOH, phenol conc. 10 wt%, at 78°C), 0.65 (neat, at 60°C) and 2.5 (in CCl₄, phenol conc. 5 wt%, at 78°C). A selective *para*-chlorination of phenol has been achieved by using CuCl₂ in DMF.⁶¹⁾ In the regioselective *para*-chlorination of *o*-cresol with SO₂Cl₂, the addition of diphenyl sulfide and AlCl₃ is effective for an improvement of the selectivity.⁶³⁾ In the chlorination of anisole with Cl₂, the *o*:*p* ratio is known to be 0.25 in CCl₄, but the use of nitromethane as solvent appears to increase the *para*-selectivity.⁶²⁾ Recently, the chlorination of phenols with good regioselectivity is achieved by using a suitable polychlorohexadienone. Phenol reacts with 2,3,4,4,5,6-hexachlorocyclohexa-2,5-dien-1-one to give *p*-chlorophenol predominantly, and with 2,3,4,5,6,6-hexachlorocyclohexa-2,4-dien-1-one to afford *o*-chlorophenol preferentially. The reaction appears to proceed *via* a charge transfer complex in which phenol acts as both π -donor and proton donor and the dienone as π -acceptor and proton acceptor.⁶⁴⁾ Several data on selective halogenation of phenols and anilines are shown in Table X.

The halogenation of moderately active and less active aromatics with molecular halogens is mostly carried out in the presence of a catalyst. However, the use of

Table X. Selective Halogenation of Phenols and Anilines.

Substrate	Halogenating Agent	Solv. and Temp. (°C)	Products (Total yields (%) & their ratio)	Ref.
PhOH	I ₂ -CuCl ₂	PhCl, 130	<i>p</i> -IC ₆ H ₄ OH (69)	56
PhNMe ₂	I ₂ -CuCl ₂	C ₆ H ₆ , 60	<i>p</i> -IC ₆ H ₄ NMe ₂ (47)	56
PhOH	I ₂ -TiOAc	AcOH, 20	<i>o</i> -IC ₆ H ₄ OH (60)	57
	Br ₂	CCl ₄ , 25	<i>o</i> - & <i>p</i> -BrC ₆ H ₄ OH (>95, 11:89)	58
	NBS ^{a)}	CCl ₄ , 25	<i>o</i> - & <i>p</i> -BrC ₆ H ₄ OH (>95, 86:14)	58
	NBS	MeCN, 25	<i>o</i> - & <i>p</i> -BrC ₆ H ₄ OH (>95, 3:97)	58
	NBS	DMF, 25	<i>p</i> -BrC ₆ H ₄ OH (70)	59
<i>m</i> -MeC ₆ H ₄ OH	NBS	DMF, 25	4-Br-3-MeC ₆ H ₃ OH (89)	59
PhNH ₂	NBS	DMF, 25	<i>p</i> -BrC ₆ H ₄ NH ₂ (93)	59
<i>p</i> -MeC ₆ H ₄ OH	Br ₂	HF-SbF ₅ , 45	3-Br-4-MeC ₆ H ₃ OH (85)	60
3,4-Me ₂ C ₆ H ₃ OH	Br ₂	HF-SbF ₅ , 45	5-Br-3,4-Me ₂ C ₆ H ₂ OH (83)	60
PhOH	CuCl ₂ -LiCl	DMF, 150	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ OH (60, 1~2:8~9)	61
PhOMe	Cl ₂	MeNO ₂ -CH ₂ Cl ₂ , 0	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ OMe (96, 1:9)	62
<i>o</i> -MeC ₆ H ₄ OH	SO ₂ Cl ₂	neat, 25	2- & 4-Cl-6-MeC ₆ H ₃ OH (97, 1:6.2)	63
	SO ₂ Cl ₂ -Ph ₂ S-AlCl ₃	neat, 15	2- & 4-Cl-6-MeC ₆ H ₃ OH (99, 1:19)	63
PhOH	<i>p</i> -hexachlorodienone ^{b)}	DMF, 20	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ OH (~100, 27:73)	64
	<i>o</i> -hexachlorodienone ^{c)}	CCl ₄ , -5	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ OH (~100, 92:8)	64

a) N-Bromosuccinimide.

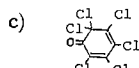
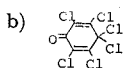


Table XI. Selective Halogenation of Aromatics with Metal Halides.

Substrate	Metal Halides	Solv. and Temp. (°C)	Products (Total yield (%) & their ratio)	Ref.
PhBr	I ₂ -SbCl ₅	CCl ₄ , 76	<i>p</i> -IC ₆ H ₄ Br (73)	65
PhF	I ₂ -SbCl ₅	CCl ₄ , 76	<i>o</i> - & <i>p</i> -IC ₆ H ₄ F (74, 2:98)	65
PhCO ₂ Et	I ₂ -SbCl ₅	ClCH ₂ CH ₂ Cl, 84	<i>m</i> -IC ₆ H ₄ CO ₂ Et (48)	65
PhBr	Br ₂ -SbCl ₅	CCl ₄ , 76	<i>o</i> - & <i>p</i> -BrC ₆ H ₄ (86, 5:95)	65
PhCl	Br ₂ -SbCl ₅	CCl ₄ , 76	<i>o</i> - & <i>p</i> -BrC ₆ H ₄ Cl (84, 4:96)	65
PhNO ₂	Br ₂ -SbCl ₅	ClCH ₂ CH ₂ Cl, 84	<i>m</i> -BrC ₆ H ₄ NO ₂ (63)	65
PhMe	FeCl ₃	neat, 40	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ Me (71, 1:5)	66
	FeCl ₃ -MoCl ₄	neat, 50	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ Me (85, 1:12)	66
	FeCl ₃ -TiCl ₄	neat, 50	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ Me (83, 1:10.5)	66
	FeCl ₃	neat, 115	<i>o</i> - & <i>p</i> -Cl ₂ C ₆ H ₄ (94, 1:12)	66
PhCl	FeCl ₃ -MoCl ₄	neat, 115	<i>o</i> - & <i>p</i> -Cl ₂ C ₆ H ₄ (86, 1:19.5)	66
	FeCl ₃ -TiCl ₄	neat, 105	<i>o</i> - & <i>p</i> -Cl ₂ C ₆ H ₄ (99, 1:20.5)	66
PhMe	TiCl ₄ -CF ₃ CO ₃ H	CH ₂ Cl ₂ , 20	<i>o</i> - & <i>p</i> -ClC ₆ H ₄ Me (90, 85:15)	67
PhPh	TiCl ₄ -CF ₃ CO ₃ H	CH ₂ Cl ₂ , 20	2- & 4-ClC ₆ H ₄ Ph (76, 80:20)	67

metal halides (involving interhalogens formed *in situ* from metal halides) frequently leads to a significant improvement of the regioselectivity and/or yield of the products. Some recent data on the selective halogenation of aromatics by means of metal halides are collected in Table XI. Equimolar mixtures of I₂ or Br₂ and SbCl₅ are found to be elegant reagents for the iodination or bromination of less reactive aromatics such as benzoic ester and nitrobenzene. Further, *p*-iodo- or *p*-bromohalogenobenzenes can be prepared by these reagents with high selectivities.⁶⁵⁾ In the chlorination of toluene with Cl₂, the *o*:*p* ratio varies with solvent considerably; for example, the ratio is 1.5 in AcOH and 0.52 in CH₃NO₂. When FeCl₃ is used as a chlorinating agent without solvent, the ratio decreases to 0.2. Recently, in the chlorination of toluene and chlorobenzene with FeCl₃, the addition of MoCl₄ or TiCl₄ is found to result in a remarkable increase in the proportion of *p*-substituted products.⁶⁶⁾ On the contrary, for toluene and biphenyl, TiCl₄ can act as an *ortho*-selective halogenating agent in the presence of trifluoroperacetic acid.⁶⁷⁾ In this case, the true reagent appears to be HOCl, though the observed *o*:*p* ratio (5.7) for toluene is somewhat higher than that (3.2) in the HOCl chlorination.

Compared to Cl₂, F₂ is much more reactive and there appears to be less selective as electrophile. This is recently confirmed by examining both orientation and relative rate in the F₂ fluorination of aromatics (in CFCl₃ at -78°C).⁶⁸⁾ For introduction of SCN into moderately active aromatics, *in situ* BrSCN or ClSCN reagent, which is more reactive than (SCN)₂, is used in the presence of a Lewis acid.^{69,70)} In the thiocyanation of alkylbenzenes and halogenobenzenes with a mixture of Pb(SCN)₂ and SbCl₅ (equivalent to *in situ* ClSCN and SbCl₅), predominant *para*-substitution is observed (for example, the *o*:*p* ratio for ethylbenzene is 0.36 in CCl₄).⁷⁰⁾ The selenocyanation of aniline is achieved by the use of Cu(OAc)₂-NaSeCN mixture and only the *para*-substitution product is isolated, but the procedure is not applicable to other active aromatics such as phenol and anisole.⁷¹⁾

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