




# SOLVENT SELECTION

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# IMPORTANCE OF SOLVENTS

- Reaction medium (transport, combine, separate)
  - Dissolution
  - (In)solubility
  - Kinetics
  - Health
  - Safety
  - Environmental aspects
  - Course of the reaction
  - Cost (purchase, recycle, dispose)
- 

# PERSPECTIVE ON SOLVENT SELECTION

ASPECT	COMMENT
Safety	Avoid solvents that are toxic or highly flammable
Promote high-yielding reactions	Compatible with desired chemistry; Can isolate product in good yield?
Convenient (minimize processing operations)	Isolate product from reaction solvent? Operate at high concentrations?
Water miscibility	Azeotroping ability Control amount of water
Cost of bulk and recoverability	More important at the end of development cycle
Environmental	Ethics, and cost of recovery and non-compliance
Long-term availability	
Acceptability for human use	
Water as solvent	But, recovery of product from aqueous layer can be costly, plus cost of disposal
Neat reactions	
Ionic liquids	


# SOLVENT SELECTION GUIDES

Almost every company created specific solvent guide  
 SmithKline Beecham – Curzons, A.D. *et al Clean Products and Processes 1, 82 (1999)*

SOLVENT		Waste	Impact	Health	Safety
Alcohols	Ethylene glycol	4	9	8	10
	1-Butanol	5	7	8	8
	Diethylene glycol mono butyl ether	5	8	8	10
	Ethanol / IMS	3	7	9	6
	2-Propanol	3	10	7	7
	Methanol	3	8	4	8
	2-Methoxy ethanol	4	9	2	7
Esters	Butyl acetate	7	7	7	6
	Propyl acetate	7	6	7	6
	Isopropyl acetate	5	7	7	6
	Ethyl acetate	4	9	7	4
	Methyl acetate	2	6	5	5
Aromatics	Xylene	8	4	5	5
	Toluene	7	3	5	4

# SOLVENT SELECTION GUIDES

Sanofi – Prat D. *et al Org.Process Res. Dev.* 17, 1517 (2013)

Solvents Guide	ETHERS: OVERVIEW					 SANOFI
Name	Overall ranking	ICH limit (ppm)	Occ. health	Safety	Environment	Other concern
<a href="#">Diethyl ether</a>	Banned	5000	OEBV2	SHB5	EHB2	Peroxides, VOC
<a href="#">Diisopropyl ether</a>	Substitution advisable	Not listed	OEBV2	SHB5	EHB3	Peroxides
<a href="#">Dibutyl ether</a>	Substitution advisable	Not listed	OEBV2	SHB5	EHB3	Peroxides, odor
<a href="#">THF</a>	Substitution advisable	720	OEBV3 Sk	SHB4	EHB2	VOC, miscible with water, peroxides
<a href="#">Methyl-THF</a>	Recommended	Not listed	OEBV2	SHB4	EHB3	Peroxides, cost
<a href="#">Dioxane</a>	Substitution requested	380	OEBV3 Sk	SHB5	EHB2	Miscible with water, peroxides
<a href="#">Anisole</a>	Recommended	5000	OEBV2	SHB3	EHB2	Odor
<a href="#">MTBE</a>	Substitution advisable	5000	OEBV3 Sk	SHB5	EHB3	VOC
<a href="#">ETBE</a>	Substitution requested	Not listed	OEBV4	SHB5	EHB3	Peroxides, lack of data
<a href="#">CPME</a>	Substitution requested	Not listed	OEBV3	SHB5	EHB3	Peroxides, one supplier only
<a href="#">Dimethoxy ethane</a>	Substitution requested	100	OEBV4 G2	SHB4	EHB2	CMR (R1B), peroxides
<a href="#">Diglyme</a>	Substitution requested	Not listed	OEBV4 G2	SHB4	EHB2	CMR (R1B), peroxides
<a href="#">Diethoxymethane</a>	Substitution requested	Not listed	OEBV4	SHB5	Not available	Reactive, considered as CMR

# SOLVENT SELECTION GUIDES

GlaxoSmithKline – Henderson, R.K. *et al Green Chemistry* 13, 854 (2011)

Solvent	Cas number	Melting point °C	Boiling Point °C	Waste	Environmental Impact	Health	Flammability & Explosio	Reactivity/ Stability	Life Cycle Score	Legislation Flag
Isopropyl acetate	108-21-4	-73	89	5	7	7	6	9	7	
Dimethyl carbonate	616-38-6	-1	91	4	8	7	6	10	8	
Ethyl acetate	141-78-6	-84	77	4	8	8	4	8	6	
t-Butylmethyl ether	1634-04-4	-109	55	4	5	5	3	9	8	
2-Methyltetrahydrofuran	96-47-9	-137	78	4	5	4	3	6	4	
Dichloromethane	75-09-2	-95	40	3	6	4	6	9	7	
Chloroform	67-66-3	-64	61	3	6	3	6	9	6	



# SOLVENT SELECTION GUIDES

ACS Green Chemistry Institute® Roundtable Solvent Selection Guide, March 2011

Substance Information			Scoring Information				
Solvent Class	Solvent Name	CAS Number	Safety	Health	Env (Air)	Env (Water)	Env (Waste)
Acid	ACETIC ACID	64-19-7	3	6	6	3	6
Acid	ACETIC ANHYDRIDE	108-24-7	3	6	6	2	7
Acid	FORMIC ACID	64-18-6	2	6	5	4	7
Acid	METHANE SULPHONIC ACID	75-75-2			6	6	10
Acid	PROPIONIC ACID	79-09-4	2	5	6	4	6
Alcohol	1-BUTANOL	71-36-3	3	5	5	5	3
Alcohol	1-PROPANOL	71-23-8	4	4	6	2	6
Alcohol	2-BUTANOL	78-92-2	4	5	6	3	5
Alcohol	2-METHOXYETHANOL	109-86-4	4	9	5	3	7
Alcohol	BENZYL ALCOHOL	100-51-6	4	3	4	2	4
Alcohol	ETHANOL	64-17-5	4	3	5	1	6
Alcohol	ETHYLENE GLYCOL	107-21-1	3	3	5	1	7
Alcohol	ISOAMYL ALCOHOL	123-51-3	3	4	5	3	4
Alcohol	ISOBUTANOL	78-83-1	3	5	4	3	3
Alcohol	ISOPROPYL ALCOHOL (IPA)	67-63-0	5	5	6	2	6
Alcohol	METHANOL	67-56-1	3	5	6	3	6
Alcohol	T-BUTANOL	75-65-0	3	5	7	2	6

# SOLVENT CONSIDERATIONS

Watch out hydrocarbon solvents with even number of carbons (toxicity, electrostatic buildup);

Classification of solvents – ICH Harmonised Guideline Q3C – Impurities: Guideline for Residual Solvents

Class 1 – solvents to be avoided (known human carcinogens, strongly suspected human carcinogens, and/or environmental hazards, e.g. carbon tetrachloride (concentration limit 4 ppm), 1,2-dichloroethane (5 ppm), 1,1,1-trichloroethane (1500 ppm), benzene (2 ppm))

Class 2 – solvents to be limited (non-genotoxic animal carcinogens, agents of irreversible toxicity, e.g. acetonitrile (410 ppm), chlorobenzene (360 ppm), chloroform (60 ppm), *N,N*-dimethylformamide (880 ppm), hexane (290 ppm), methanol (3000 ppm), *N*-methylpyrrolidone (530 ppm), toluene (890 ppm))

Class 3 – solvents with low toxic potential (permissible daily exposure 50 mg or more per day, e.g. acetic acid, acetone, ethyl acetate, heptane, 2-propanol, triethylamine)

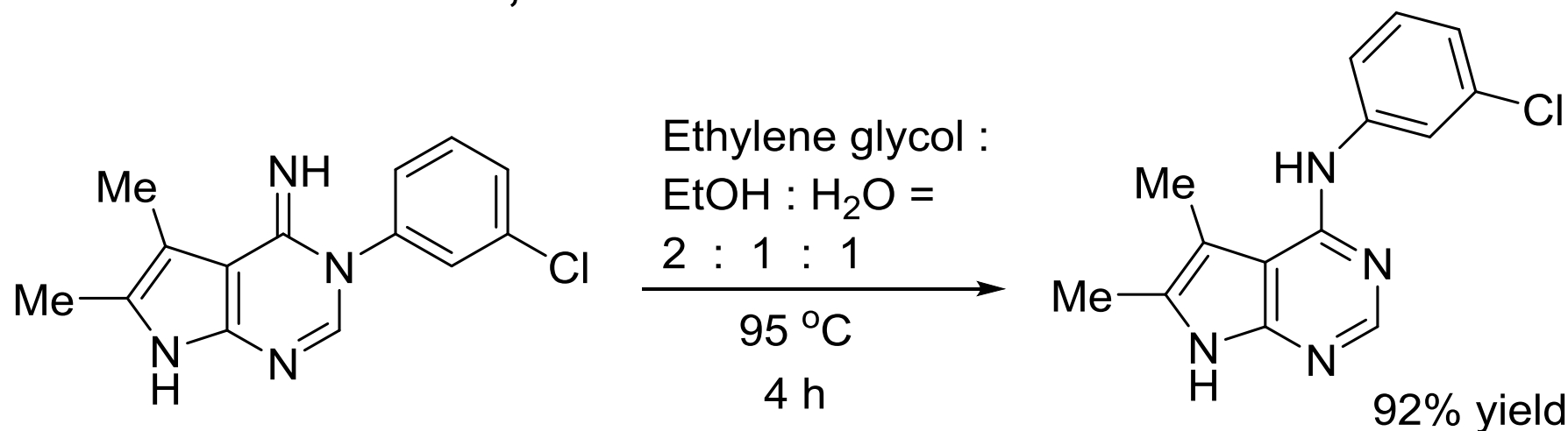
Solvents for which no adequate toxicological data was found – a manufacturer is asked to supply justification for residual levels of these solvents (e.g. diisopropyl ether, petroleum ether, trifluoroacetic acid)



# SOLVENT CONSIDERATIONS

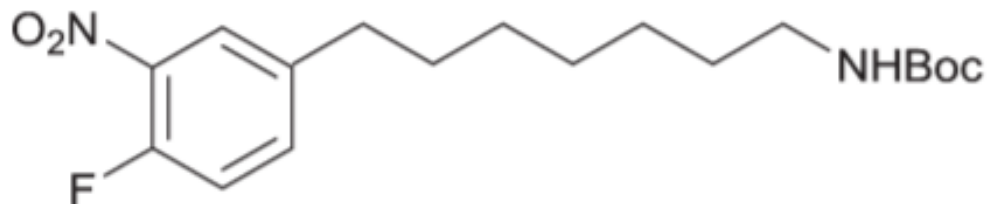
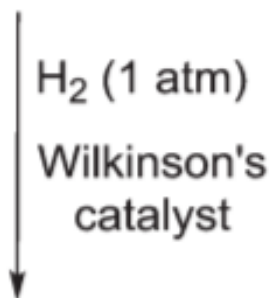
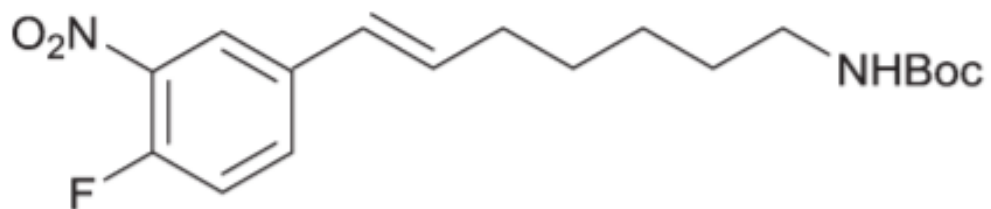
The best reaction solvent is the one that crystallizes the product directly from the reaction;

Novartis – the Dimroth rearrangement – temperature and solubility turned out to be the most important – the product simply precipitated from the reaction mixture;



Fischer, R.W. *Org. Process Res. Dev.* 5, 581 (2001)

# SOLVENT CONSIDERATIONS



Solvent	Yield (% after 48 h)
Benzene	0
Toluene	20
Methanol	80
THF	91
Methanol-THF	93

Dyson, P.J.; Jessop, P.G. *Catal. Sci. Technol.* **6**, 3302 (2016)



# SOLVENT CONSIDERATIONS

Homogeneous vs. Heterogeneous  
Reactions using gases

Insolubility is sometimes advantageous (the Schotten-Baumann reaction, the Finkelstein reaction)

Menshutkin (1890)

The reaction rate of the reaction of triethylamine with alkyl halides providing quaternary ammonium salts strongly depends on a solvent (hexane **1**, acetone **338**, benzyl alcohol **739**)

Solvents commonly used in **academia** are not often welcomed for **industrial** applications

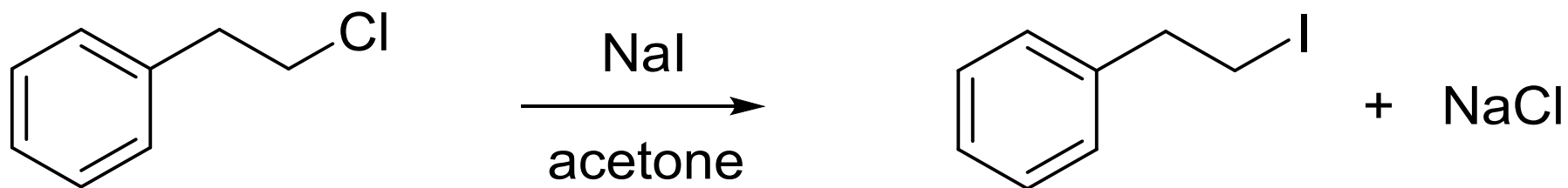
Safety first !



# SOLVENT CONSIDERATIONS

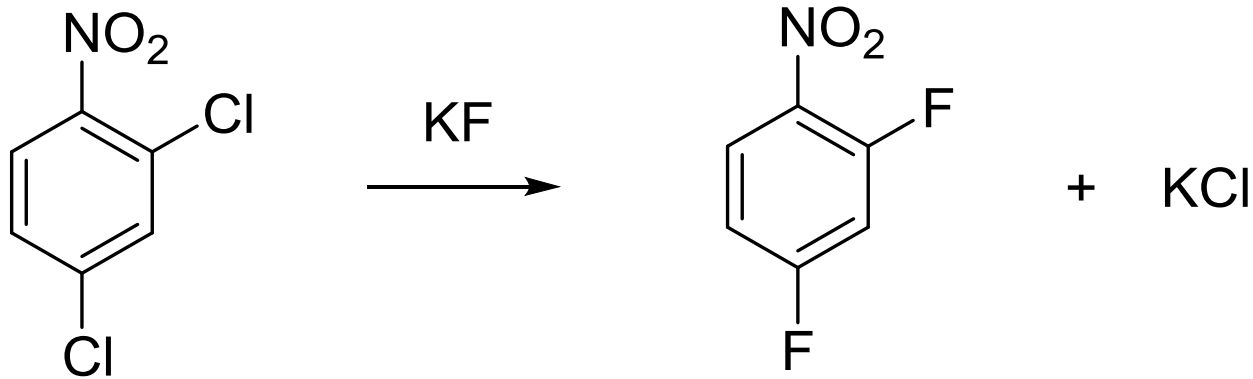
## The Finkelstein reaction

The reaction driven to completion by exploiting the different solubility of used and formed halides



# SOLVENT CONSIDERATIONS

## The Halex reaction

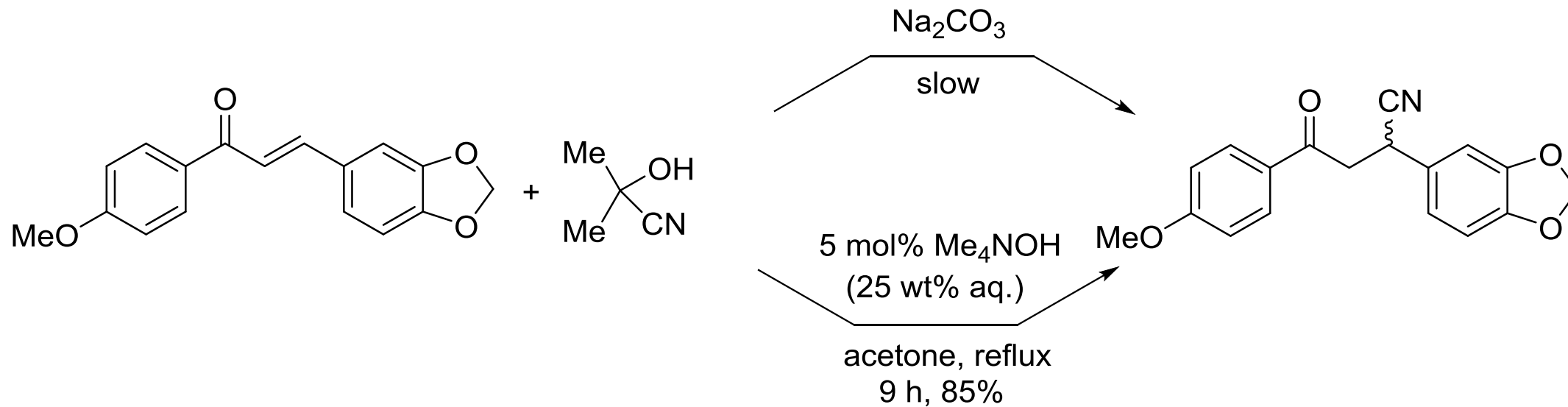


toluene ..... no reaction

water ..... no reaction

DMF ..... satisfying conversion, KCl precipitates out of the reaction mixture

# MIXING OF HETEROGENEOUS MIXTURES



Anderson, N.G. *Practical Process Research and Development, A guide for organic chemists*, 2nd Edition, Elsevier 2000



# SOLVENT CONSIDERATIONS


Solvent	TWA (ppm)
Acetone	500
EtOAc	400
MeOH	200
<i>t</i> -BuOH	100
MTBE	50
MeCN	20
DMF	10
Pyridine	1
2-Methoxyethanol	0.1

TWA = Time-Weighted Average shift for five days for nearly safe exposure over an 8 h shift for five days for nearly all workers



# SOLVENT CONSIDERATIONS

Also, always consider physical-chemical properties like

- Flash point
  - Flammability
  - Boiling point
  - Melting point
  - Electrostatic charge accumulation
  - Recycling potential
  - Cost of solvent
  - Environmental aspects
  - Cost of disposal
  - Polarity
- 

# SOLVENT CONSIDERATIONS

Solvents **rarely used** in the pharmaceutical industry

Solvent	Disadvantage	Alternative replacement
Diethylether	Flammable	MTBE
Diisopropylether	Peroxide formation	MTBE
Hexane	Electrostatic charge Neurological toxicity	Heptanes, <i>i</i> -octane
Chloroform	Mutagenicity, environmental aspects, toxicity	Dichloromethane, 2-MeTHF, toluene
Benzene	Toxicity	Toluene
Ethylene glycol	Toxicity	1,2-Propandiol
Acetonitrile	Animal teratogen, potential acetamide generation (genotoxic)	2-propanol, acetone - water

# SOLVENT CONSIDERATIONS

Solvents **preferred** for process development (Pfizer)

Preferred	Usable	Undesirable
Water	Cyclohexane	Pentane
Acetone	Heptane	Hexanes
Ethanol	Toluene	Diisopropyl ether
2-Propanol	Methyl cyclohexane	Diethyl ether
Ethyl acetate	MTBE	Dichloroethane
<i>i</i> -Propyl acetate	<i>i</i> -Octane	<b>Dichloromethane</b>
Methanol	2-MeTHF	Chloroform
Methyl ethyl ketone	DMSO	DMF
<i>n</i> -Butanol	AcOH	NMP
<i>t</i> -Butanol	Ethylene glycol	1,4-Dioxane
		Benzene
		Carbon tetrachloride



# SOLVENT CONSIDERATIONS



## Polarity of solvents

**Hughes-Ingold rules** ((de)stabilization of transition state)

Aliphatic nucleophilic substitution and elimination reactions;  
Considering **pure electrostatic interactions** between ions or dipolar molecules and solvent molecules in **initial** and **transition** states;



# SOLVENT CONSIDERATIONS

## Hughes-Ingold rules

Reaction type	Initial reactants	Activated complex	Charge alteration during activation	Effect of increased solvent polarity on rate <sup>a)</sup>
(a) S <sub>N</sub> 1	R—X	R <sup>δ+</sup> ... X <sup>δ-</sup>	Separation of unlike charges	Large increase
(b) S <sub>N</sub> 1	R—X <sup>+</sup>	R <sup>δ+</sup> ... X <sup>δ+</sup>	Dispersal of charge	Small decrease
(c) S <sub>N</sub> 2	Y + R—X	Y <sup>δ+</sup> ... R ... X <sup>δ-</sup>	Separation of unlike charges	Large increase
(d) S <sub>N</sub> 2	Y <sup>-</sup> + R—X	Y <sup>δ-</sup> ... R ... X <sup>δ-</sup>	Dispersal of charge	Small decrease
(e) S <sub>N</sub> 2	Y + R—X <sup>+</sup>	Y <sup>δ+</sup> ... R ... X <sup>δ+</sup>	Dispersal of charge	Small decrease
(f) S <sub>N</sub> 2	Y <sup>-</sup> + R—X <sup>+</sup>	Y <sup>δ-</sup> ... R ... X <sup>δ+</sup>	Destruction of charge	Large decrease



# SOLVENT CONSIDERATIONS

## Hughes-Ingold rules

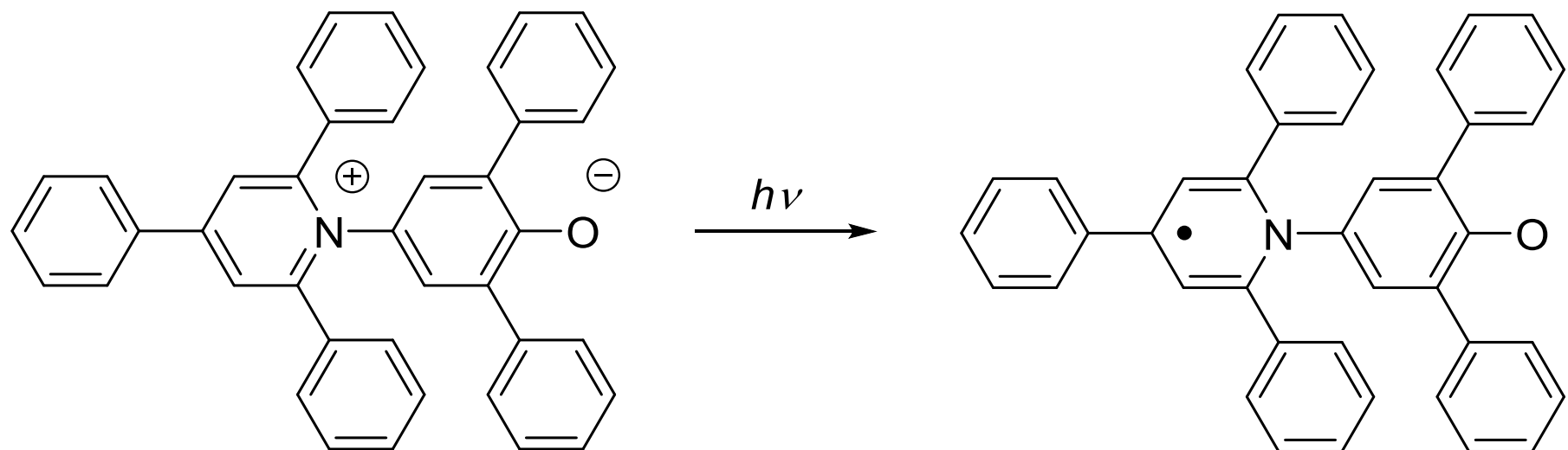
- An **increase** in solvent **polarity** results in an **increase** in the **rates** of those reactions in which the charge density is **greater** in the activated complex than in the initial reactant molecules;
- An **increase** in solvent **polarity** results in a **decrease** in the **rates** of those reactions in which the charge density is **lower** than in the initial reactant molecules;
- A change in solvent polarity will have **negligible effect** on the **rates** of those reactions that involve **little** or **no** change in the charge density from reactants to the activated complex.

# SOLVENT CONSIDERATIONS

## Polarity of solvents

Reichardt, C. *Pure Appl. Chem* 76, 1903 (2004)

Reichardt, C. *Solvents and Solvent Effects in Organic Chemistry*,  
3 rd Ed., Wiley-VCH, 2003



# SOLVENT CONSIDERATIONS

## Polarity of solvents

$$E_T(30) = h \cdot c \cdot N_A \cdot \nu_{max} \quad [\text{kcal} \cdot \text{mol}^{-1}]$$

Could be used even in solid rare gas matrices to establish polarity of rare gases (Xe → Kr → Ar → Ne decreasing polarizability)

$E_T(30)$  recommended and used for monitoring of a quality of disinfection solutions of alcohol/water in order to prevent SARS-CoV 2 infections

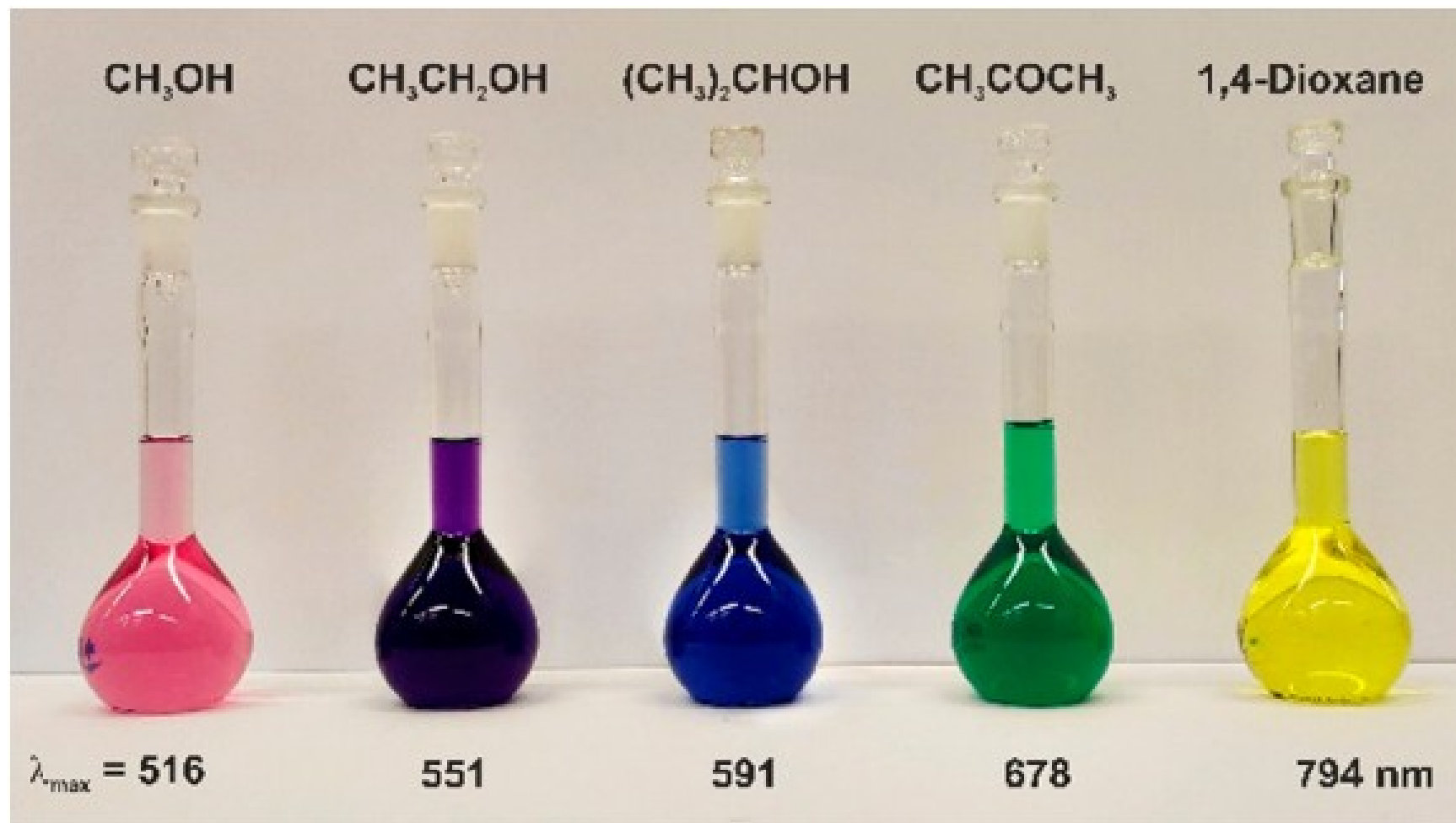
# SOLVENT CONSIDERATIONS

## Polarity of solvents

$E_T^N$  parameter – the negative solvatochromism of the  $\pi \rightarrow \pi^*$  shifts of solutions of the betaine dye – more polar solvents stabilize the ground energy of the polar dye, producing thus greater shift in the position of  $\pi \rightarrow \pi^*$  absorption relative to that found for solutions of the dye in tetramethylsilane.

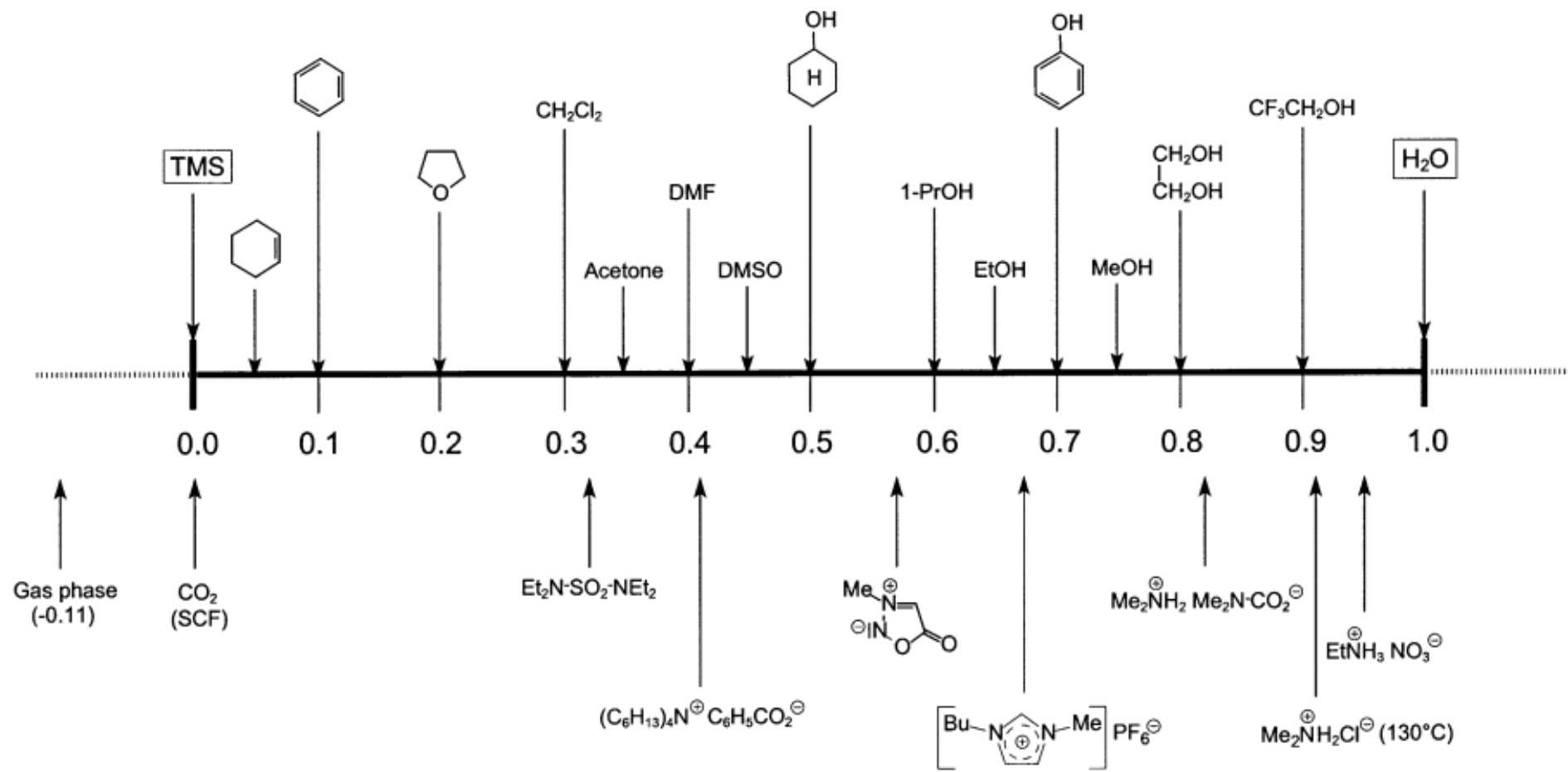
Colors of this dye in a solvent are indicative of the polarities of the solvent and solvent combination used to dissolve it.

# SOLVENT CONSIDERATIONS



Reichardt, C. *J.Org.Chem* 87, 1616 (2022)

# SOLVENT CONSIDERATIONS



Reichardt, C. *Pure Appl. Chem.* 76, 1903 (2004)



# SOLVENT CONSIDERATIONS

## Polarity of solvents

Solvent	Polarity $E_T^N$	Solubility in water (wt%)	Bp of water-solvent azeotrope	wt% of water removed by azeotrope	ICH solvent class
Water	1.000	-	None	None	
EtOH	0.654	$\infty$	78 °C	4.0	3
AcOH	0.648	$\infty$	77 °C	97	3
DMF	0.404	$\infty$	None	None	2
Acetone	0.355	$\infty$	None	None	3
CH <sub>2</sub> Cl <sub>2</sub>	0.309	1.3	38 °C	1.5	2
Toluene	0.099	0.06	84 °C	13.5	2
Et <sub>3</sub> N	0.043	5.5	75 °C	10	3
Heptane(s)	0.012	0.0004	79 °C	12.9	3
Cyclohexane	0.006	0.006	69 °C	9	2

# SOLVENT CONSIDERATIONS

## Polarity of solvent mixtures

Solvent	Polarity $E_T^N$	Solvent mixture	Calculated $E_T^N$
MeOH	0.762	EtOH:H <sub>2</sub> O = 6.9:3.1	0.762
EtOH	0.654	Acetone:H <sub>2</sub> O = 4.6:5.4	0.654
H <sub>2</sub> O:CH <sub>2</sub> Cl <sub>2</sub> = 0.2:99.8	0.310	H <sub>2</sub> O:MIBK = 1.9:98.1	0.283
H <sub>2</sub> O:CH <sub>2</sub> Cl <sub>2</sub> = 0.2:99.8	0.310	H <sub>2</sub> O:EtOAc = 3.3:96.7	0.253
H <sub>2</sub> O:CH <sub>2</sub> Cl <sub>2</sub> = 0.2:99.8	0.310	H <sub>2</sub> O:2-MeTHF = 5.3:94.7	0.223
<i>i</i> -PrOAc	0.210	Heptanes:EtOAc = 0.8:9.2	0.210
MeOH:H <sub>2</sub> O = 7:1	0.792	EtOH:H <sub>2</sub> O = 5:3	0.783

# SOLVENT CONSIDERATIONS

- Tendency of solvents to form **azeotropes with water** is considered advantageous (it is not practical and economical to dry solvents using drying agents on large scale);
- **Be careful** – dependence on pressure (**breaking the azeotrope**)

Effect of reducing distillation pressure on EtOAc – water:

Pressure (mm)	Bp (°C)	Water in azeotrope (wt%)
760	70.4	8.5
250	42.6	6.3
25	1.9	3.6

# IMPURITIES IN SOLVENTS

- Absolute solvents are rather expensive, common solvents contains some amount of water, for certain operations they should be dried (azeotropic distillation, molecular sieves, use of an excess of cheap reagent);

Denatured solvents (ethanol)

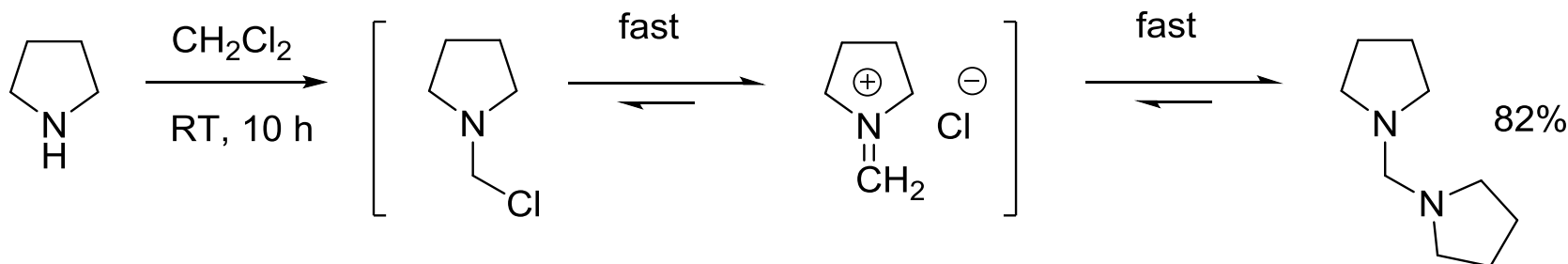
Stabilizers (e.g. BHT in THF)

Tendency to form (hydrogen)peroxides (diisopropyl ether, butadiene, acetaldehyde, 1,4-dioxane, styrene, acrylonitrile, 2-butanol, benzyl alcohol, THF, MIBK, 2-propyl alcohol)

# IMPURITIES IN SOLVENTS

- Degradation of solvents (ethyl acetate, DMF)

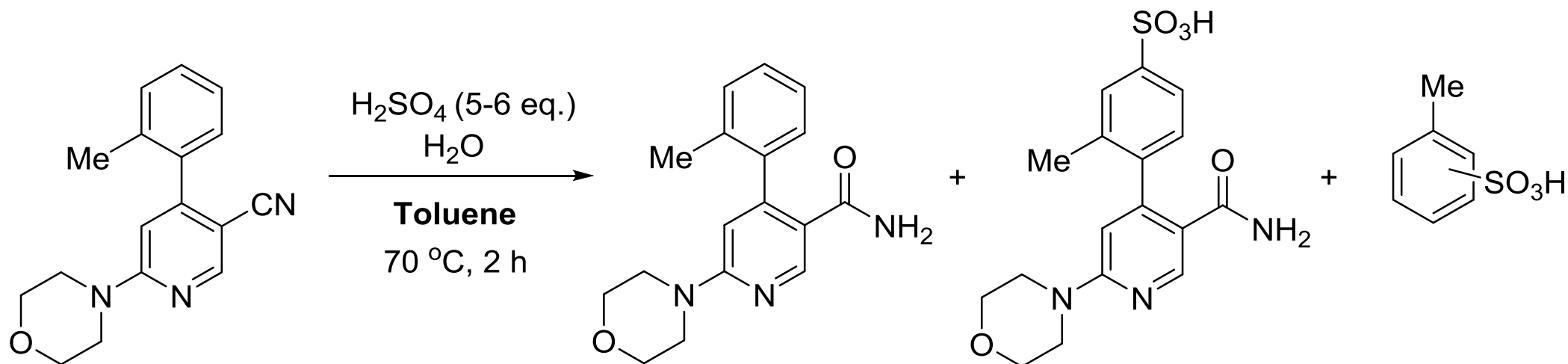
Side reactions (reesterifications, reaction with dichloromethane)



Avoid unwanted formation of esters of sulfonic acids (potentially mutagenic)

# SACRIFICIAL SOLVENTS

About 50 eq. of 98%  $\text{H}_2\text{SO}_4$  at 50 °C for 3 h followed by an aqueous quench provided ring sulfonation in the product; 5-6 eq. of 98%  $\text{H}_2\text{SO}_4$  in toluene at 70 °C for 2 h – sulfonation of the product was significantly diminished.

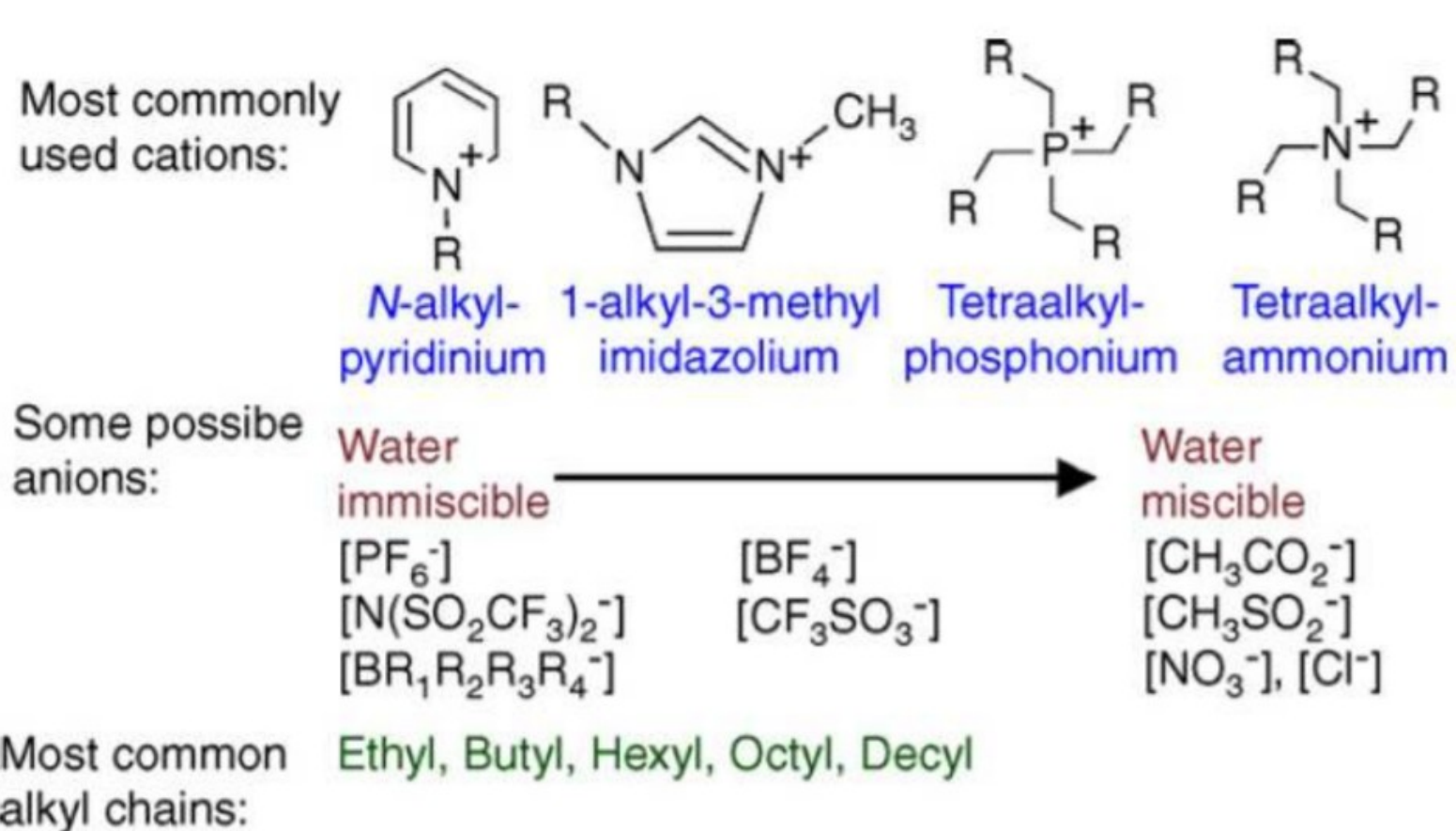


Harrington, P.J. *et al* *Org. Process Res. Dev.* 10, 1157 (2006)

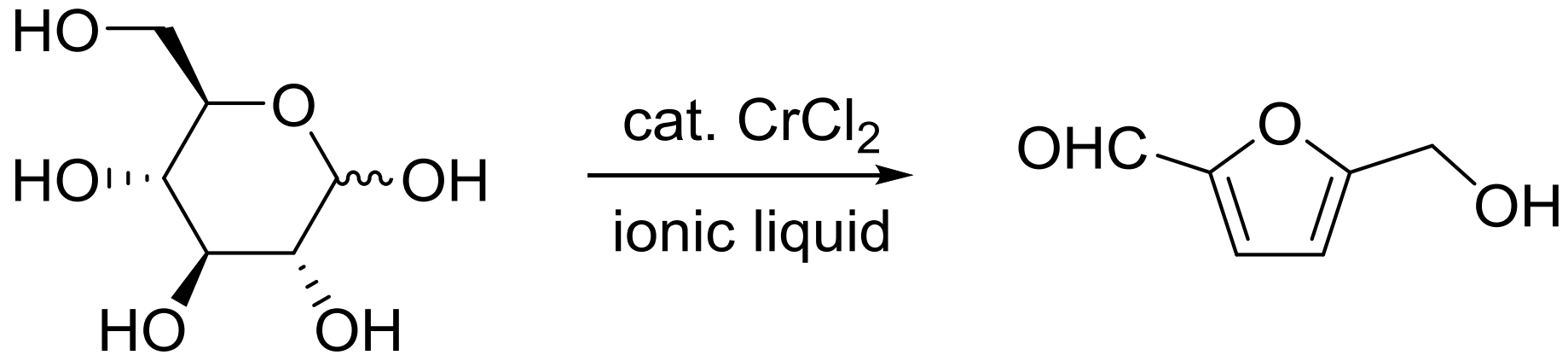
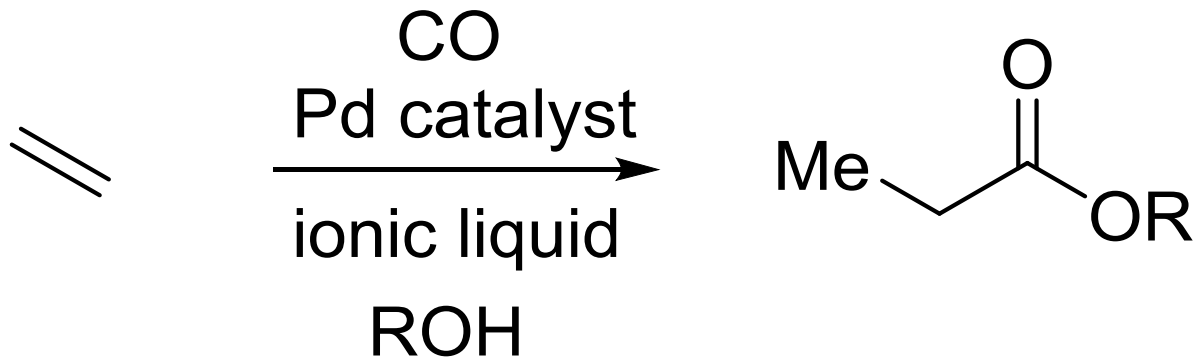


# IONIC LIQUIDS

- Ionic liquids are ionic compounds (salts) which are liquids below 100 °C. More commonly, ionic liquids have melting points below room temperature.



# IONIC LIQUIDS

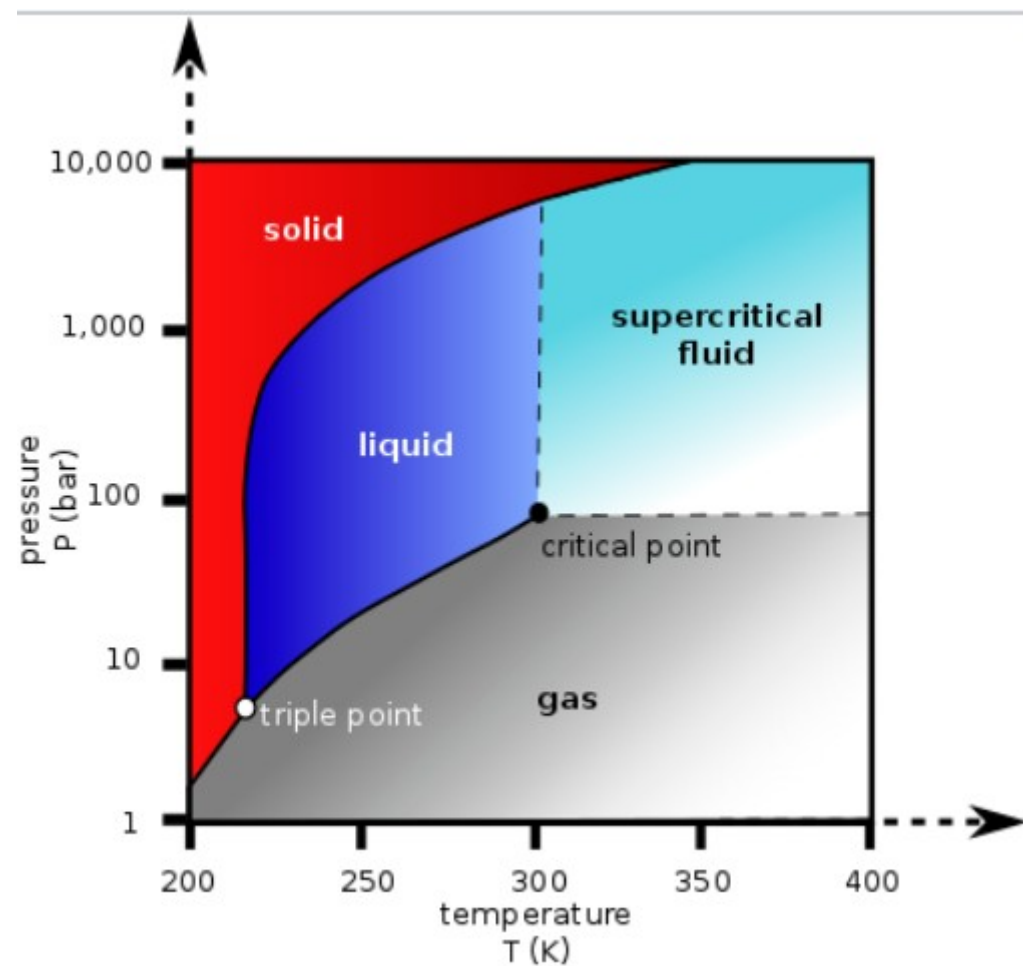


# SUPERCRITICAL FLUIDS

SCF	Name	$T_c$ (°C)	$p_c$ (bar)	$d_c$ (g/mL)	MW	$\mu$ (De- bye) <sup>a</sup>	Cost <sup>b</sup> (\$/kg)
Ar	argon	-122.5	48.6	0.531	39.95	0	6
CO <sub>2</sub>	carbon dioxide	31.1	73.8	0.466	44.01	0	3
HCl	hydrogen chloride	51.5	82.6	0.42	36.46	1.08	20
HBr	hydrogen bromide	90.0	85.5	n.a.	80.91	0.82	50
HI	hydrogen iodide	150.7	83	n.a.	127.9	0.44	n.a.
H <sub>2</sub> O	water	374.0	220.6	0.322	18.02	1.85	n.a.
NH <sub>3</sub>	ammonia	132.4	113.2	0.235	17.03	1.47	3
N <sub>2</sub> O	nitrous oxide	36.4	72.5	0.453	44.01	0.167	50
Kr	krypton	-63.76	54.9	0.912	83.80	0	3000
SF <sub>6</sub>	sulfur hexafluoride	45.5	37.6	0.737	146.1	0	50
Xe	xenon	16.6	58.3	1.099	131.3	0	4000

Jessop, P.G.; Leitner, W. *Chemical Synthesis Using Supercritical Fluids*, Wiley-VCH 1999

# SUPERCRITICAL CARBON DIOXIDE



Peach, J.; Eastoe, J. *Beilstein J.Org.Chem.* 10, 1878 (2014)

Beckman, E.J. *J.Supercritical Fluids* 28, 121 (2004)



# **SUPERCRITICAL CARBON DIOXIDE**

## **FOOD INDUSTRY**

Coffee decaffeination

Tea decaffeination

Fatty acids from spent barley

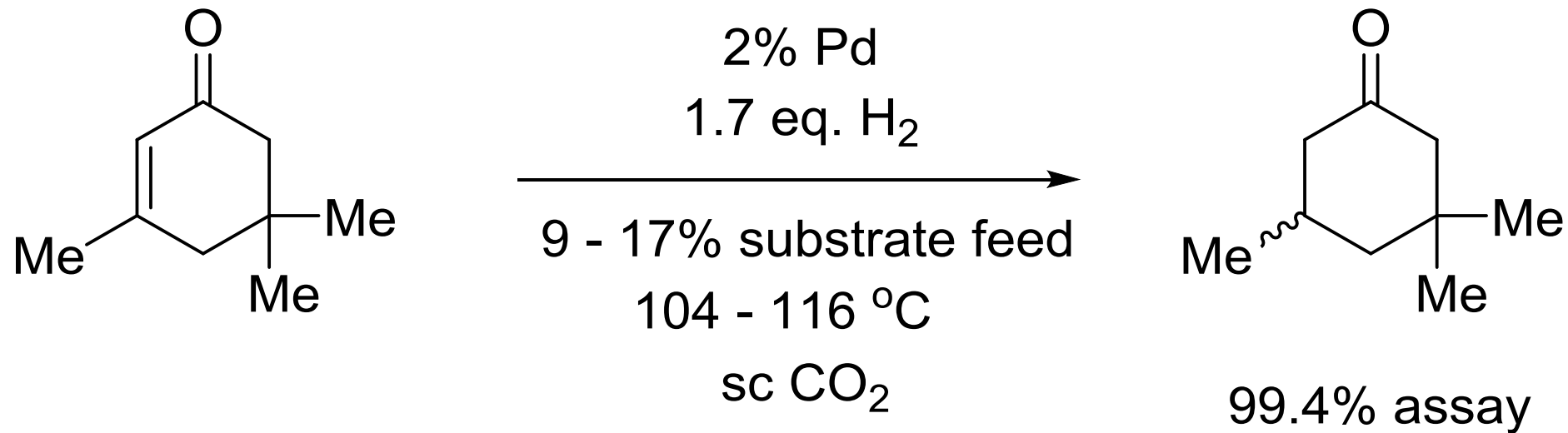
Vitamin E, hops, spices extraction

Nicotine extraction

Natural insecticide/pesticide extraction



# SUPERCRITICAL CARBON DIOXIDE



Thomas Swan & Co., Ltd.

Licence, P. *et al* *Green Chem.* 5, 99 (2003)



## GAS EXPANDED LIQUIDS (GXLs)

Combination of a classical solvent and supercritical solvent (usually  $\text{scCO}_2$ ) – adjustable polarity;

