Annexure III Syllabi and course structure of M.Tech Degree Course In Industrial Catalysis

SEMESTER I

Course code	Paper	Credit	C/E	E Marks		Total
				Tests and Assignments	End Sem Exam	marks
CHE 3101	Surface Chemistry and Catalysis	4	C	60	40	100
CHE 3102	Catalyst Technology – I	4	C	60	40	100
CHE 3103	Physical Methods In Catalysis	4	С	60	40	100
CHE 3104	Chemical Reaction Engineering	3	E	60	40	100
CHE 3105	Enzyme Catalysis	3	Е	60	40	100
CHE 3106	Electrocatalysis	3	Е	60	40	100
CHE 3107	Lab course	2	С	100		100
CHE 3108	Viva voce (End Semester)		C		50	50

SEMESTER II

Course code	Paper	Credit	C/E	Marks		Total
				Tests and Assig- nments	End Sem	marks
CHE 3201	Surface Characterization Techniques	4	С	60	40	100
CHE 3202	Catalysis by Metal Complexes	4	С	60	40	100
CHE 3203	Catalyst Technology – II	4	С	60	40	100
CHE 3204	Industrial Catalytic Processes	3	Е	60	40	100
CHE 3205	Phase Transfer Catalysis	3	Е	60	40	100
CHE 3206	Polymer Supported Catalysis	3	Е	60	40	100
CHE 3207	Lab course	2	С	100		100
CHE 3208	Viva voce (End Semester)		С		50	50

SEMESTER III

Course code	Paper	Credits	C/E	Total marks
CHE 3301	Project progress evaluation	16	С	300

SEMESTER IV

Course code	Paper	Credits	C/E	Total marks
CHE 3401	Project Dissertation evaluation &	16	С	300
	Viva voce			

SEMESTER I

CHE 3101 SURFACE CHEMISTRY& CATALYSIS

4 credits

64 hours

UNIT I

The solid - liquid interface – surface energy from solubility changes – surface energy from immersion, – contact angle – contact angle hysterisis – experimental methods and measurement of contact angle – theories of contact angle phenomena – adsorption of non-electrolytes from dilute solutions – irreversible adsorption – adsorption in binary liquid systems – adsorption of electrolytes.

UNIT II

The solid - gas interface – surface area of solids – structure and chemical nature of solid surfaces – nature of the solid adsorbate complex – adsorption of gases and vapours on solids – adsorption time – Langmuir adsorption isotherm – the BET and related isotherms – isotherms based on equation of state of the adsorbed films – potential theory – phase transformations in the multiplayer region – thermodynamics of adsorption – critical comparison of various models for adsorption.

UNIT III

Physical adsorption on heterogeneous surfaces – rate of adsorption – adsorption on porous solids: hysterisis – chemisorption and catalysis – chemisorption isotherms – kinetics of chemisorption – surface mobility – chemisorption bond – heterogeneous catalysis – transition state theory – specificity and selectivity in catalysis – catalytic activity and strength of chemisorption – Langmuir-Hinshelwood and Eley-Rideal mechanism – electronic factors in catalysis by metals – geometric factors in heterogeneous catalysis.

UNIT IV

Chemical dynamics at surfaces – inelastic collision and trapping – the sticking coefficient – physisorbed molecules – molecule migration, desorption and transition to chemisorbed state – accommodation coefficients – energy exchange at surface – cube models. Precursor theory of chemisorption – sticking probability curves – their temperature dependence. Desorption – analysis of desorption curves – kinetics of adsorption and desorption.

UNIT V

Dynamics of selective catalysis– catalyst selectivity –kinetic method of estimating selectivity in consecutive reactions – localized and mobile adsorption of the intermediate product – effect of pore size on catalyst selectivity – poly-functional catalysis –inter-particle space and through gas phase – simplified criteria for negligible resistance to mass transfer – selectivity of poly-functional catalysis.

- 1. A.W. Adamson & A.P. Gast, "Physical Chemistry of Surfaces", John Wiley and Sons, Canada, 1997.
- 2. R.H.P. Gasser, "An introduction to chemisorption and catalysis by metals", Oxford, 1985.
- 3. A. Clark, "Theory of adsorption and catalysis", Academic Press, New York, 1970.
- 4. D.K. Chakraborthy, "Adsorption and catalysis by solids", Wiley Eastern Ltd. 1990.
- 5. J.M. Thomas & W.J. Thomas, "Introduction to principles of heterogeneous catalysis", Academic Press, New York, 1967.
- 6. G. A. Somorjai, "Introduction to Surface Chemistry and Catalysis", Wiley-Black Well, New York, 1994

CHE 3102 CATALYST TECHNOLOGY – I

64 hours

4 credits

UNIT I

Catalyst preparation methods – precipitation and coprecipitation – sol gel process–Dispersed metal catalysts; support materials; preparation and structure of supports; surface properties – preparation of catalysts – introduction of precursor compound – interaction of metal compound with substrate surface – metal distribution within catalyst pellets – metal cluster compounds as active precursors – pre-activation treatment – drying and calcinations – activation process.

UNIT II

Bulk catalysts and supports – fused catalysts – Skeletal metal catalysts.Heteropoly compounds – Solid superacids. Carbons – structural chemistry of carbon– basic structures – loosely defined structures – formation of carbon materials: catalytic formation of carbon from molecules – carbon on noble metal catalysts – carbon formation in zeolites – graphitization of carbon – reaction of oxygen with carbon – surface chemistry of carbon – non-oxygen hetero elements on carbon surfaces – surface oxygen groups – carbon as catalyst support – carbon as catalyst.

UNIT III

Synthesis of aluminosilicate zeolites and related silica based materials – structure, composition, zeolite synthesis, mechanism and chemistry–zeolites obtained from various reaction systems – synthesis of some selected important zeolites – titanosilicates – activation of zeolites. Modification of zeolites – ion exchange – metals supported on zeolites – reduction of metal ions in zeolite – dealumination of zeolites; Shape selective catalysis in zeolites; Phosphate based zeolites and molecular sieves.

UNIT IV

Pillared clays– properties of pillared clays–synthesis and catalytic applications of pillared clays– the use of coordination and organometallic compounds as pillaring agents – the use of polymers and surfactants in pillaring process – pillaring of acid activated clays – enhancement of acidity of PILCs – the use of PILCs as catalyst supports –hydrotalcite like anionic clays in catalytic organic reactions; Mesoporous materials - ordered mesoporous materials – PMOS -synthesis of silica molecular sieve materials - characterization of mesoporous molecular sieves – catalytic properties of mesoporous materials.

UNIT V

Catalyst manufacture– scope and goals – catalysts prepared by precipitation – solution and slurry transfer – filtration – drying: calcining; ion exchange; pulverization, pilling and extrusion; crushing and screening to produce granules; coating (not impregnation); impregnation to orient the coating material to the support – anchor coating or wash coating.

- 1. G. Ertl, H. Knozinger and J. Weitkamp (eds), "Preparation of Solid Catalysts", Wiley-VCH, Verlag, 1999.
- 2. J.R. Anderson and M. Boudart (Eds), "Catalysis, Science and Technology", Vol 6, Springer-Verlag, BerlinHeildberg, 1984.
- 3. J. Weitkamp and L. Puppe (eds), "Catalysis and zeolites fundamentals and applications", Springer-Verlag Berlin Heidelberg 1999.
- 4. A. Corma, Chemical Reviews, 97 (1997) 2373-2419.

- 5. S. van Donk, A.H. Janssen, J.H. Bitter & K.P. deJong, Catalysis Reviews, 45 (2003) 297-319.
- 6. A. Gil, L.M. Gandia& M.A. Vincente, Catalysis Reviews Science and Engineering, 42 (2000) 145-212.
- 7. B.F. Sels, D.E. De Vos & P.A. Jacobs, CatalysisReviews, 43 (2001) 443-488.
- 8. M. Hartmann & L. Kevan, Chemical Reviews, 99 (1999) 635-663.
- 9. A.B. Stiles and T.A. Koch, "Catalyst manufacture", Marcel Dekker Inc., NY, 1995.

CHE 3103 PHYSICAL METHODS IN CATALYSIS

64 hours

Determination of surface area and pore structure of catalysts – physical adsorption methods – mercury intrusion methods – chemisorption methods – X-ray methods – microscopic methods – radioactive isotope method – flow rate and diffusion methods – miscellaneous methods – comparison of methods; Measurement of acidity of surfaces – aqueous methods – nonaqueous liquid phase methods – adsorption from gas phase – hydrogen-deuterium exchange reactions.

UNIT II

Gas chromatography – basic instrumental setup – carriers – columns – detectors – scientific selection of stationary phases – applications in catalytic activity studies – determination of surface area of catalysts – principles and instrumentation of Gas Chromatography–Mass Spectrometry (GC-MS); TG, DTA, DSC – Instrumentation – application in catalysis.

UNIT III

Conventional magnetic methods in catalysis – forms of magnetism – force produced by magnetism – techniques – applications. UV-Vis-NIR spectroscopy – general background – electronic transitions – molecular complexes – non molecular solids – vibrational transitions – DRS – theoretical background – experimental considerations – applications.

UNIT IV

Infrared spectroscopy – experimental techniques – application of infrared spectroscopy to adsorption and catalysis; Raman spectroscopy – theory of Raman Effect – application of Raman spectroscopy to catalysis; Mossbauer spectroscopy – occurrence of the Mossbauer effect – method of measurement – hyperfine interactions – isomer effect – quadrupole splitting – magnetic splitting – applications to adsorption and catalysis.

UNIT V

Electron spin resonance spectroscopy – features of ESR spectra – g value – shifts in g value – tensor properties of g and A parameter – origin of hyperfine interaction – Fermi contact and dipolar terms – identification of catalytic sites by EPR – detection of free radical intermediates – molecular motion on the surface.

Nuclear magnetic resonance spectroscopy – solid state NMR – basic principles and methods of solid state NMR – fundamentals of NMR spectroscopy – nuclear spin interactions in solids – dipolar interaction – chemical shift interaction – quadrupolar interaction – experimental techniques – dipolar decoupling – MAS – DOR – CP – 2D NMR - useful nuclei – important NMR parameters and related structural information – applications.

REFERENCE

- 1. R.B. Anderson, "Experimental methods in catalysis research", Vol I, II, Academic press, New York, 1981.
- 2. W.M.A. Niessen, "Current Practice of Gas Chromatography Mass Spectrometry", Marcel Dekker Inc, New York, 2001.
- 3. W.N. Delgass, G.L. Haller, R. Kellerman and J.H. Lunsford, "Spectroscopy in heterogeneous catalysis", Academic press, New York, 1979.
- 4. G. Ertl, H. Knozinger and J. Weitkamp, "Handbook of Heterogeneous Catalysis" Vol 2, Wiley-VCH, Weinheim, 1997.

CHE 3104(E) CHEMICAL REACTION ENGINEERING

48 hours

Overview of chemical reaction engineering – homogeneous reactions in ideal reactors – kinetics of homogeneous reactions – concentration dependent term – temperature dependent term – searching for a mechanism – predictability of reaction rate from theory – interpretation of batch reactor data – constant volume and varying volume batch reactor – temperature and reaction rate.

UNIT II

Introduction to reactor design – ideal reactors for a single reaction – ideal batch reactor – steady state mixed flow reactor – steady state plug flow reactor; Design for single reactions – size comparison of single reactors – multiple reactor systems – recycle reactor – autocatalytic reactions; design for parallel reactions.

UNIT III

Potpourri of multiple reactions – irreversible first order reactions in series – first order followed by zero order – zero order followed by first order – successive irreversible reactions of different orders – reversible reactions – irreversible series-parallel reactions; Temperature and pressure effects – single reactions – multiple reactions – choosing the right kind of reactor.

UNIT IV

Reactions catalyzed by solids – heterogeneous reactions – solid catalyzed reactions – rate equation for surface kinetics – pore diffusion resistance combined with surface kinetics – porous catalyst particles – heat effects during reaction – performance equations for reactors containing porous catalyst particles – experimental methods for finding rates – product distribution in multiple reactions; the packed bed catalytic reactor; Reactors with suspended solid catalyst-fluidized reactors of various types (mention only); G/L reactions on solid catalyst: trickle beds, slurry reactors, three-phase fluidized beds (mention only).

UNIT V

Biochemical reaction systems – enzyme fermentation – Michaelis-Menten kinetics – inhibition by a foreign substance; Microbial fermentation – introduction and overall picture; substrate-limiting microbial fermentation – batch (or plug-flow) fermentors – mixed flow fermentors – optimum operations of fermentors; product limiting microbial fermentation – batch or plug-flow fermentors – mixed flow fermentors.

REFERENCE

- 1. O. Levinspeil, "Chemical Reaction Engineering", IIIrd edition, John Wiley and sons, New York, 1999.
- 2. J.M. Smith, "Chemical Engineering Kinetics", IIIrd edition, McGraw Hill Inc. Singapore, 1981.
- 3. H.S. Fogler, "Elements of Chemical Reaction Engineering", IInd edition, Prentice Hall of India Pvt. Ltd. New Delhi, 1997.

CHE 3105 (E) ENZYME CATALYSIS

UNIT I

Enzymes - classification and nomenclature of enzymes – structure of enzymes – effect on reaction rate – thermodynamic definitions – catalytic power and specificity of enzymes – optimization of weak interactions between enzyme and substrate in the transition state – binding energy, reaction specificity and catalysis – specific catalytic groups contributing to catalysis – enzyme kinetics as an approach to understanding mechanism – substrate concentration and rate of enzyme catalyzed reactions – quantitative relationship between substrate concentration and reaction rate – kinetic parameters used to compare enzyme activities.

UNIT II

Immobilized biocatalysts – definition and classification of immobilized biocatalysts – reasons for immobilization – history of immobilization – economic importance; methods for immobilization – adsorption – ionic binding – covalent binding – cross-linking – matrix entrapment – membrane confinement – combined methods – immobilization on crosslinked polymer supports- immobilization of coenzymes. Synthesis of peptides using immobilized enzymes- triphasic catalysis.

UNIT III

Characteristics of immobilized biocatalysts – activity as a function of temperature – stability as a function of temperature – temperature optimum in the long term process – the influence of pH value – influence of substrate concentration – influence of diffusion – other physical properties; Reactors for immobilized biocatalysts – stirred reactors – loop reactors – bed reactors – membrane reactors – special forms of reactors.

UNIT IV

Lewis acid properties of Zn and its development to phosphotriester detoxifying agents – vanadium haloperoxidases – Mo and W enzymes – catalysis by nitrogenases and synthetic analogs – biological Fe-S clusters with catalytic activity.

UNIT V

Catalysis by Ni in biological systems – oxygen activation at non-heme iron centers – dioxygen activation at heme centers in enzymes and synthetic analogs – biological and biomimetic catalysis of Mn redox enzymes and their inorganic models.

REFERENCE

- 1. A.L. Lehninger, "Principles of Biochemistry", Worth Publishers, USA, 1987.
- 2. W. Hartmeier, "Immobilized Biocatalysts: an introduction", Springer-Verlag, BerlinHeidelberg, 1986.
- 3. J. Reedijk and E. Bouwman, "Bioinorganic Catalysis", Marcel Dekker Inc. New York, 1999.

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48 hours

CHE 3106 (E) ELECTROCATALYSIS

UNIT I

Hydrogen adsorption at metal surfaces- importance of surface cleanliness –principles of hydrogen-metal interactions – classification of different cases of hydrogen adsorption – physisorption on metal surfaces – activated and non-activated adsorption of hydrogen on metal surfaces – hydrogen chemisorption including surface restructuring – subsurface, dissolved and hydridic hydrogen; hydrogen on Pt systems as example of importance of hydrogen-metal interaction in catalysis and electrochemistry.

UNIT II

The science of electrocatalysis on bimetallic surfaces – bimetallic surface chemistry – bulk alloys – overlayers and surface alloys– oxidation of adsorbed CO – oxidation of dissolved CO – formic acid oxidation – oxidation of methanol; fundamental aspects of vacuum and electrocatalytic reactions of methanol and formic acid on Pt surfaces.Fuel Cells- different types of Fuel Cells - fuel cell efficiency and technical anode potential – reaction rates by electrical measurements – non-electrical measures of reaction rates – techniques for mechanistic studies – UHV models of electrocatalysis.

UNIT III

Electrocatalytic hydrogenation of organic compounds – reaction mechanism – ECH on noble metals – ECH at electrodes made of Raney type materials; Recent advances in kinetics of oxygen reduction – gas phase studies of oxygen- surface structure and reaction kinetics – surfaces modified by foreign metal adatoms – non-noble metals and metal oxides – transition metal and macrocyclic complexes – electrolyte effects – temperature dependence of transfer coefficient for oxygen reduction.

UNIT IV

Studying electrocatalytic oxidations of small organic molecules with in-situ IR spectroscopy – principles and methods of in-situ IR spectroscopy – dissociative adsorption of small organic molecules at the surface of Pt group metals – molecular structure effects in electrocatalytic oxidation of small organic molecules – surface structure effects in electrocatalytic oxidation of small organic molecules – in-situ time resolved IR spectroscopic studies of the kinetics of electrocatalytic oxidation of small organic molecules.

UNIT V

Unified model of electron and ion transfer reactions at metal electrodes – extended Anderson-Newns model – activation barrier and transfer coefficient – pre-exponential factor – double layer effects in the electrode kinetics of electron and ion transfer reactions– electro-reductions of anions – double layer effects at single crystal electrodes – ion transfer reactions – electron transfer to adsorbed reactants – electron and proton transfer at self assembled monolayer.

REFERENCE

- 1. J. Lipkowski and P.N. Ross, "Electrocatalysis", Wiley VCH, Canada, 1998.
- 2. J. O'M. Bockris and S.U.M. Khan, "Surface Electrochemistry", Plenum press, New York, 1993.
- 3. J. O'M. Bockris, A.K.N. Reddy and M. G-. Aldeco, "Modern Electrochemistry fundamentals of electrodics", IInd edition, Kluwer academic/Plenum press, New York, 2000.

CHE 3107 PRACTICAL – I

9 hours / week

3 credits

- 1. Preparation of supports for catalysts alumina, silica, amorphous silica-alumina, zeolites, chromia, titania.
- 2. Comparison of different preparation techniques precipitation, precipitation from homogeneous solution, polymer pyrrolysis, template synthesis, sol-gel method etc.
- 3. Shaping of support materials to extrudates and spheres.
- 4. Preparation of dispersed metal oxide catalysts co precipitation, impregnation techniques.
- 5. Conversion of active phase into metal form reduction.
- 6. Determination of surface area of the catalysts using BET method, total and micro pore volume and evaluation of pore size and particle size distribution.
- 7. Thermal studies of catalysts using TG and DTA/DSC.
- 8. Measurement of acidity of catalysts Temperature Programmed Desorption of ammonia, adsorbed ammonia IR spectra, TG of pyridine, 2,6 DMP adsorbed samples.
- 9. Characterization of surface functional groups using IR spectroscopy.
- 10. Fabrication of reactors.

SEMESTER II

CHE 3201 SURFACE CHARACTERIZATION TECHNIQUES

4 credits

64 hours

UNIT I

Surface crystallography and electron diffraction – surface symmetry – description of over-layer structures – reciprocal net and electron diffraction – surface structure determination using low energy electron diffraction (LEED)– reflection high energy electron diffraction (RHEED) – powder XRD – principles of powder diffraction – phase identification – quantitative analysis – determination of atomic structure of crystalline catalysts – determination of local atomic arrangement of amorphous catalysts – extended X-ray absorption fine structure spectroscopy (EXAFS).

UNIT II

X-ray photoelectron spectroscopy (XPS) – photon sources – shapes and shifts – XPS as core level spectroscopy – synchrotron radiation studies – structural effects in XPS – Auger electron spectroscopy (AES) – basic processes – energy levels – shifts and shapes – AES for surface composition analysis – comparison of AES and XPS. Ultraviolet photoelectron spectroscopy (UPS) – UPS in the study of adsorbed molecules.

UNIT III

Incident ion techniques – charge exchange between ions and surfaces – applications in ion scattering techniques – low energy ion scattering (LEIS) – application of LEIS in catalysis – secondary neutral mass spectrometry (SIMS) - secondary ion mass spectrometry (SIMS) – theory of SIMS – electron and photon emission under ion bombardment – energy distribution of secondary ions – ionization probability – emission of molecular clusters – conditions for static SIMS – charging of insulating samples – applications in catalysts – model catalysts.

UNIT IV

Desorption spectroscopies – thermal desorption techniques – qualitative analysis of pressuretime curves – experimental arrangements for flash desorption and TPD, FD and TPD spectra – temperature programmed reduction – thermodynamics of reduction – reduction mechanisms – applications; temperature programmed sulphidation; electronically stimulated desorption – basic mechanisms, instrumentation and applications.

UNIT V

High field techniques – field emission – field emission microscope – factors governing operation – practical microscope configurations – applications of field emission microscopy (FEM);Scanning tunneling and atomic force microscopy (STM, AFM); Work function techniques – single crystal surfaces – polycrystalline surfaces – work function measurements based on diode method – field emission measurements – photoelectric measurements; Electron microscopy (SEM, TEM) – instrumentation – applications on catalysts – element analysis in the electron microscope.

- 1. D.P. Woodruff and T.A. Delchar, "Modern techniques of surface science", CambridgeUniversity press, 1990.
- 2. J.W. Niemantsverdriet, "Spectroscopy in Catalysis: an introduction", VCH, NY, 1995.
- 3. G. Ertl, H. Knozinger and J. Weitkamp, "Handbook of Heterogeneous Catalysis" Vol 2 & 3, Wiley-VCH, Weinheim, 1997.

CHE 3202 CATALYSIS BY METAL COMPLEXES

4 credits

64 hours

UNIT I

Coordinative unsaturation– Oxidative addition- Addition reactions of hydrogen, halogens, organic halides- Addition reactions of Si-H and C-C bonds- C-H activation – Reductive elimination reactions- Migration (insertion reaction)- Insertion of CO, isocyanide, carbon dioxide, alkene and alkynes- nucleophilic and electrophilic attack on coordinated ligands- Ligand cone angles– catalytic cycles – hard and soft catalysis.

UNIT II

Hydrogenation reactions- Reversible cis-dihydrido catalysts- asymmetric hydrogenation- alkene isomerisation, hydrosilylation and hydroboration reaction- Reactions of Co and hydrogen- watergas shift reaction- Fischer –Tropsch reaction- hydroformylation of unsaturated compounds carbonylation reactions- Oxo synthesis – production of acetic acid by carbonylation of methanol – selective ethylene oxidation by the Wacker process – oxidation of cyclohexane- Monsanto L-Dopa process

UNIT III

Oligomerization of ethylene (SHOP); Cluster compounds as homogeneous catalysts and catalyst precursors – classification of catalyst precursors based on structure. homogeneous and heterogeneous Zeigler-Natta catalysts – supported metal complex catalysts – Phillips process for ethylene polymerization; Late-metal catalysts for ethylene homo and copolymerization – effect of nature of metallocene complexes of group IV metals on their performance in catalytic ethylene and propylene polymerization – carbocationic alkene polymerizations initiated by organo transition metal complexes.

UNIT IV

Polymer bound transition metal complex catalysts – synthesis of supports and catalysts – characterization by physical methods – catalysis – stability of polymer supported catalysts – comparison of polymers with inorganic catalyst supports; Specific features of catalysis by immobilized metal complexes – ligand exchange in metal-polymeric systems – characteristics of electron-transfer reactions – macromolecular effects – main factors regulating activity – effects of cluster formation and cooperative stabilization – outlook for polyfunctional catalysis – technological aspects.

UNIT V

Zeolite entrapped metal complexes – synthesis – flexible ligand method – ship in a bottle method – zeolite synthesis method – characterization – stability analysis – oxygen adsorption – cyclovoltametry – catalysis by zeolite entrapped transition metal complexes – ordered mesoporous and microporous molecular sieves functionalized with transition metal complexes as catalysts for selective organic transformations – propene polymerization with silica supported metallocene/MAO catalysts.

- 1. F.A. Cotton, G. Wilkinson, C.A. Murillo & M. Bochmann, "Advanced inorganic chemistry", VI edition, John Wiley and sons, Singapore, 1999.
- 2. J.E. Huheey, "Inorganic chemistry: principles of structure and reactivity", Harper and Row publishers, Singapore.

- 3. J. Hagen, "Industrial catalysis: A practical approach", Wiley VCH, Weinheim, Germany, 1999.
- 4. D.C. Sherrington & P. Hodge, "Synthesis and separations using functional polymers", John Wiley and Sons, 1988.
- 5. A.D. Pomogailo, "Catalysis by polymer-immobilized metal complexes", Gordon & Breach Science Publishers, Amsterdam, 1998.
- 6. G. Ertl, H. Knozinger and J. Weitkamp, "Handbook of Heterogeneous Catalysis" Vol 3 and 5, Wiley-VCH, Weinheim, 1997.

CHE 3203 CATALYST TECHNOLOGY – II

4 credits

64 hours

UNIT I

Deactivation of catalysts – classification of catalyst deactivation processes; general aspects of catalyst deactivation – poisoning of catalysts – poisoning of metallic catalysts – poisoning of non metallic catalysts – poisoning of bifunctional catalysts – coke formation on catalysts – metal deposition on catalysts – sintering of catalysts; diffusion and deactivation of catalysts – analogy between selectivity and deactivation mechanisms – catalyst deactivation – correlations for activity decay – separable and non separable kinetics; Optimization of deactivated reactor systems – comparison of various reactor types under deactivating conditions – optimization of deactivating reactors – optimal temperature policies – other optimal policies.

UNIT II

Regeneration of deactivated catalysts – feasibility of regeneration – description of coke deposit and kinetics of regeneration – regeneration of fluidized bed reactors – regeneration of coked catalyst pellets – regeneration of fixed beds containing coked catalysts; Technological economics – cost of producing a chemical – variable costs – fixed costs – direct costs – indirect charges – capital dependent charges – effect of scale of operation – effect of low plant operation – contribution of catalyst to production cost; Catalytic processes in an integrated system – effect of catalyst on plant equipment – catalyst life – effect of improvement in catalyst performance on process and economics – prospects for new catalytic process

UNIT III

Design of industrial catalysts; design procedure; the overall design of catalysts – overall development of an industrial catalyst – scientific basis of design – the idea – preliminary checking – the description of the idea – theoretical design: primary components of the catalyst – catalyst deactivation – secondary components of a catalyst – selection of the preferred form of a catalyst – the overall design; design of the primary constituent of the catalyst – theories of chemical bonding – theories of bonding and adsorption – bond theories and catalysis – theories of bonds and catalyst design – activity patterns and catalyst design.

UNIT IV

Design of the secondary components of a catalyst – secondary component design via mechanistic studies – alloy catalysts – metal cluster catalysts – metal oxide solid solutions – specific examples of oxide solid solution catalysts applied to design; choice of support materials – texture and strength of support – chemical interaction – deactivation; experimental testing – preliminary testing – testing of the effect of secondary components – measurement of reaction kinetics – tubular reactors – plug reactors – stirred reactors – long term catalyst testing.

UNIT V

Specific examples of catalyst design; design of a catalyst for conversion of olefins to aromatics – description of the idea – design of primary constituents – experimental testing – design of secondary components; design of a catalyst for selective hydrogenation of acetylene in presence of ethylene – the idea – the description of the idea – the design of the primary components; design of a catalyst for reduction of nitrogen oxides to nitrogen – the idea – description of the idea – co-ordination and geometric considerations and design – catalysis by metals – conclusions – comments.

REFERENCE

1. R. Hughes, "Deactivation of catalysts", Academic press, London, 1984.

- 2. R. Pearce and W.R. Patterson, "Catalysis and chemical processes", Academic press, Leonard Hill, London, 1981.
- C.A. Heaton (ed), "An Introduction to Industrial Chemistry", Leonard Hill, London, 1984.
 D.L. Trimm, "Design of industrial catalysts", Elsevier scientific, NY, 1980.

CHE 3204 (E) INDUSTRIAL CATALYTIC PROCESSES

3 credits

48 hours

UNIT I

Environmental catalysis – Mobilesources – Stationary sources –Catalytic routes to hydro (chloro) fluoro carbons –Inorganic reactions – Ammoniasynthesis –Ammonia oxidation.

UNIT II

Energy related catalysis – Perspectivesin oil refining – Steam reforming – Water gas shift and COS removal – Methanol synthesis – CO and CO₂ hydrogenation – Methanol to hydrocarbons – Hydrotreating reactions – Catalytic reforming – Catalytic cracking

UNIT III

Hydrocracking and catalytic dewaxing –Aromatization of light alkanes – Catalytic coal gasification – Catalysis in coal liquefaction – Fuel cells – Heterogeneous photocatalysis.

UNIT IV

Alkylation of aromatics – Isomerization and trans-isomerization – Dehydrogenation reactions – Hydrogenation reactions – Hydroformylation – Selective oxidations.Carbon-carbon bond forming reactions- coupling reactions- Ni, Cu, Pd catalysts in C-C bond forming reactions. Heck, Suzuki, Sonogashira, Stille Coupling reactions.

UNIT V

Amination reactions – Acylation of aromatics – Nucleophilic Aromatic Substitution reactions – Elimination and addition reactions – Oligomerization and metathesis.

- 1. G. Ertl, H. Knozinger and J. Weitkamp, "Handbook of Heterogeneous Catalysis" Vol 4 and 5, Wiley-VCH, Weinheim, 1997.
- 2. R.J. Farrauto and C.H. Bartholomew, "Fundamentals of Industrial Catalytic Processes", Blackie Academic and Professional Chapman and Hall, 1997.
- 3. R. Pearce and W.R. Patterson, "Catalysis and chemical processes", Academic press, Leonard Hill, London, 1981.
- 4. J. Weitkamp and L. Puppe (eds), "Catalysis and zeolites fundamentals and applications", Springer-Verlag Berlin Heidelberg 1999.

CHE 3205 (E) PHASE TRANSFER CATALYSIS

3 credits

UNIT I

Basic concepts in phase transfer catalysis – phase transfer catalyzed reactions – basic steps of phase transfer catalysis – effect of reaction variables on transfer and intrinsic rates – outline of compounds used as phase transfer catalysts; Phase transfer catalysts – use of quaternary salts – macrocyclic and macrobicyclic ligands – PEG's and related compounds – other soluble polymers – use of dual phase transfer catalyst or co-catalyst in phase transfer systems – catalysts for transfer of species other than anions – separation and recovery of phase transfer catalysts.

UNIT II

Insoluble phase transfer catalysts –PTC catalysts bound to resins – phase transfer catalysts bound to inorganic solid supports – phase transfer catalysts contained in separate liquid phase (third liquid phase catalyst); Variables in reactor design for application of PTC – choice of catalyst – structure activity relationships – catalyst stability – separation and recycle – choice of solvent and nature of chemical reaction – stabilization of transition state – solubility of complex in organic phase – rate of transfer – solvent free PTC – presence of water – agitation – choice of anion, leaving group and counter anion – choice of base – guidelines for exploring new PTC applications;

UNIT III

PTC displacement reactions with simple anions – general considerations – important factors – characteristics of various anions – PTC catalysts for displacement reactions – behaviour of various anions in PTC displacement reactions – cyanide, halide, carboxylates, azide, sulphide and disulphide, thiocyanate, sulphite, nitrite, hydroxide, carbonate and bicarbonate, peroxide and superoxide, phosphide and phosphinite, cyanate; PTC reactions with strong bases – C alkylation – N alkylation – O alkylation: etherification – S alkylation: thioetherification – dehydrohalogenation – carbene reactions – condensation reactions – deuterium exchange, isomerization etc.

UNIT IV

Phase transfer catalyzed oxidations – permanganate oxidations – oxidations with hypochlorite and hypobromite, hydrogen peroxide, air or oxygen,persulphate, nitric acid, CCl₄/NaOH, periodate and related anions,perborate, ferrate and ferricyanide, superoxide – PTC electrochemical oxidations – PTC oxidations with other oxidants;PTC reductions – NaBH₄ reductions – LiAlH₄ reductions – reductions with sodium formate, sulphur containing anions – hydrogenations – reductions with formaldehyde – electrochemical reductions – photochemical reductions – Wolff-Kishner reduction – reduction by dodecarbonyl tri iron and related species

UNIT V

PTC: Chiral phase transfer catalyzed formation of C-C bonds – alkylation reactions –methylation and alkylation – asymmetric alkylation of oxindoles – synthesis of chiral amino acids – Michael addition reactions; PTC: transition metal co-catalyzed reactions – carbonylation and reactions with CO – formation of metal carbonyl anions – carbonylation of alkyl and aryl halides, olefins, acetylenes, aziridines and azobenzenes, thiiaranes, phenol – reduction and hydrogenation with metal co-catalysts – coupling reactions of alkenes, alkynes and alkyl halides – other reactions

- 1. C.M. Starks, C.L. Liotta and M. Halpern, "Phase Transfer Catalysis fundamentals, applications and industrial perspectives", Chapman&Hall, New York, 1994.
- 2. Y. Sasson and R. Neumann (eds), "Handbook of Phase Transfer Catalysis", Chapman & Hall, Great Britan, 1997.

CHE 3206 (E) POLYMER SUPPORTED CATALYSIS

3 credits

48 hours

UNIT I

Preparation of functionalized polymers – polymerization – condensation polymerization – addition polymerization – copolymerization – chemical functionalization of synthetic organic polymers – functionalization of polystyrene – functionalization of condensation polymers – chemical modification under phase transfer catalysis – functionalization by grafting – functionalization of membranes – functionalization of biopolymers – functionalization of inorganic supports.

UNIT II

Organic reactions using polymer supported catalysts – monitoring of polymer supported reactions – some general features of polymer supported reactions – selected polymer supported catalysts – Design and industrial application of polymeric acid catalysis – homogeneous versus heterogeneous catalysis – industrial applications – new developments in catalyst design- polymer supported dendrimers and dendrigraft polymers as catalysts in organic reactions- organo catalysts.

UNIT III

Hydrogenation processes catalyzed by metal containing polymers – principal information concerning the mechanism – relationships between preparation conditions and catalytic properties – composition, structure and activity, influence of solvent properties – influence of reaction temperature – dependence of rate on concentration ratios – catalysis by dispersed colloidal particles and transition metal clusters – selectivity – enantioselective hydrogenation – catalytic hydrogenation by heterometallic polymers – H-H activation and hydrogen transfer – hydrosilylation – reduction of macromolecules.

UNIT IV

Oxidation reactions catalyzed by immobilized metal complexes – basic laws – activation and binding of dioxygen – hydrocarbon oxidation – polymeric metalloporphyrin catalyzed olefin oxidation – catalase type activity – metal polymer catalyzed olefin epoxidation by alkyl hydroperoxides – macromolecular complexes for oxidation – oxidation of sulphur containing compounds – asymmetric induction in oxidationreactions – catalytic oxidation of polymers.

UNIT V

Activation of small stable molecules and stimulation of catalytic reactions through participation of immobilized metal complexes – Fischer-Tropsch synthesis – carbonylation – water gas shift reaction – hydroformylation – catalytic hydrolysis and dehydration – acid catalysis – binding and activation of molecular hydrogen – photochemical decomposition of water – electrochemical stimulation of catalytic processes; Polymeric phase transfer catalysts – synthesis of catalysts – analysis of catalysts – mechanism of catalysis – experimental design – synthetic applications.

- 1. A. Akelah& A Moet, "Functionalized polymers and their applications", Chapman and Hall, 1990.
- 2. D.C. Sherrington & P. Hodge, "Synthesis and separations using functional polymers", John Wiley and Sons, 1988.
- 3. A.D. Pomogailo, "Catalysis by polymer-immobilized metal complexes", Gordon & Breach Science Publishers, Amsterdam, 1998.

CHE 3207 PRACTICAL – II

9 hours / week

3 credits

- 1. Phase characterization-using XRD.
- 2. Activity studies for hydrogenation reactions using pulsed micro-catalytic reactors and continuous flow reactors.
- 3. Activity studies of cracking and oxidation reactions.
- 4. Activity studies for FC Alkylation and acylation reactions.
- 5. Product analysis using Gas Chromatography.
- 6. Comparison of activity for different types of reactors STR, FBR.
- 7. Synthesis and characterization of some transition metal complex catalysts using conductance, magnetic susceptibility, IR and TG measurements.
- 8. Oxidation of p-xylene to tere-phthalic acid using Co(II)Br₂.
- 9. Hydrogenation of buta-1,3-diene using pentacyamocobaltate(II)
- 10. Oxidation of ethylene to acetaldehyde using palladium chloride.