EUROPEAN FEDERATION OF CHEMICAL ENGINEERING

WORKING PARTY ON DISTILLATION, ABSORPTION AND EXTRACTION

Recommended Systems

For

Liquid Extraction Studies

Edited by T. Mišek

Published on behalf of the European Federation of Chemical Engineering by

The Institution of Chemical Engineers, George E. Davis Building, 165-171 Railway Terrace, Rugby, Warwickshire CV21 3HQ

© 1978

All rights reserved. No part of this publication may be reproduced, stored in a retrieval system, or transmitted, in any form or by any means, electronic, mechanical, photocopying, recording or otherwise, without the prior permission of the copyright owner.

-(ISBN No. 0 85295 113 2)

Foreword

The Working Party on "Distillation, Absorption and Extraction" is one of the working parties set up by the European Federation of Chemical Engineering, a non-profit making association of technical and scientific societies which are active in the field of Chemical Engineering, Chemical Technology and Process Engineering.

The main activity of the Working Party on "Distillation, Absorption and Extraction" is directed towards organising informal discussion meetings on subjects within its terms of reference; occasionally, the Working Party also participates in large international conferences, e.g. the International Symposium on Distillation 1969 and the coming similar event of 1979.

Progress in the field of separation techniques and equipment very much depends on experimental investigations. During the Working Party's discussion meetings, it was soon concluded that the situation with respect to the use of test mixtures for determination of the performance of equipment was less than satisfactory. As a consequence, the decision was made to attempt to standardise test mixtures for distillation columns. As a result of a joint effort, the manual "Recommended Test Mixtures for Distillation Columns" was issued in 1969. This latter work mainly collates data from the literature and presents the information required for description of some ten test mixtures.

Following publication of the distillation manual, standardisation of test mixtures for liquid-liquid extraction studies was initiated. A Sub-Committee of the Working Party was set up in 1972 with Dr. T. Misek (Czechoslovakia) as the Chairman, and Dr. H.W. Brandt (West Germany) and Professor C. Hanson (Great Britain) as members.

The situation regarding the proper choice of system for liquid-liquid extraction studies was found to be even more confusing than that for distillation. Literature data on equilibria and physical properties were conflicting; thus indicating the need for experimental verification and completion of relevant physical data. After due consideration, the Sub-Committee selected three test mixtures for closer investigation. It is most gratifying that the Sub-Committee was able to secure the co-operation of Universities and Industrial laboratories for determination of the necessary data. A list of the participating individuals and organisations is presented in the introduction by the Chairman of the Sub-Committee, Dr. T. Misek. The introduction also describes the general principles upon which the work performed was based.

The preparatory work of data collection took considerably more time than originally planned and the collation of these data also proved to be time-consuming. Now that the work is complete, the Working Party is very pleased to present with this manual "Recommended Systems for Liquid Extraction Studies", the results of the study to which so many people contributed in various ways.

Special thanks have to be extended to Dr. T. Misek and the other members of the Sub-Committee for their enthusiasm and perserverance in completing this project. I would also like to gratefully acknowledge the contribution of Ing. L. Strnadova, Ing. D. Stangl, and their co-workers who assisted Dr. Misek in assembling and treating the experimental data and in the preparation of the manual.

OCTOBER 1977

Professor F.J. Zuiderweg

Chairman of the Working Party on Distillation, Absorption and Extraction

A Strategic State

PREFACE TO THE FIRST EDITION

(

The period of the last three decades has been characterized not only by rapid development in the use of liquid extraction in industry but also by associated progress in research and development of extraction equipment, processes and the behaviour of two-phase liquid systems in general.

For all these purposes, it was necessary to perform experimental work using different liquid systems the number of which is still growing. Many systems have been employed, chosen according to the specific objectives of particular studies, availability at the time or other random effects. In addition to their composition, the systems used have been defined by some of their thermodynamic and physical properties. These have served as a starting point for the quantitative description of the effects of these properties on the phenomena studied in two-phase liquid systems.

However, it has become apparent that this approach has serious drawbacks. Thus physical and thermodynamic properties chosen are not able to define completely the systems used; systematic errors and deviations are often introduced in their measurement, while formerly unknown property effects and phenomena are revealed at later stages making the research work performed with different systems uncomparable or even useless. When testing extraction equipment, the use of different systems leads necessarily to different results and to conclusions that may not be generally justified.

The recommendation of using only a few selected systems in liquid extraction studies is aimed at providing a comparable basis for these studies, and at achieving deeper knowledge of these systems, so that the above disadvantages can be avoided. Thus, in addition to testing extraction equipment in the laboratory and pilot plant, the recommended systems should also be used for studying the basic components of the extraction processes, e.g. formation and motion of drops, transfer of heat and mass, coalescence, interphase phenomena, etc. Hopefully, the work of different authors could then become comparable and the explanation of anomalous phenomena, deviations and strange behaviour will be possible.

The Sub-Committee of the Working Party faced a difficult task in selecting the systems which should be considered and eventually recommended for general use after thorough investigation. The selection necessarily involved compromises and the systems chosen for this study are not ideal. However, they can undoubtedly meet the objective and are supported by the experience of the principal institutions dealing with liquid extraction in Europe, the data in the literature and the results of the contributors obtained during this project.

PREFACE TO THE FIRST EDITION

(

The period of the last three decades has been characterized not only by rapid development in the use of liquid extraction in industry but also by associated progress in research and development of extraction equipment, processes and the behaviour of two-phase liquid systems in general.

For all these purposes, it was necessary to perform experimental work using different liquid systems the number of which is still growing. Many systems have been employed, chosen according to the specific objectives of particular studies, availability at the time or other random effects. In addition to their composition, the systems used have been defined by some of their thermodynamic and physical properties. These have served as a starting point for the quantitative description of the effects of these properties on the phenomena studied in two-phase liquid systems.

However, it has become apparent that this approach has serious drawbacks. Thus physical and thermodynamic properties chosen are not able to define completely the systems used; systematic errors and deviations are often introduced in their measurement, while formerly unknown property effects and phenomena are revealed at later stages making the research work performed with different systems uncomparable or even useless. When testing extraction equipment, the use of different systems leads necessarily to different results and to conclusions that may not be generally justified.

The recommendation of using only a few selected systems in liquid extraction studies is aimed at providing a comparable basis for these studies, and at achieving deeper knowledge of these systems, so that the above disadvantages can be avoided. Thus, in addition to testing extraction equipment in the laboratory and pilot plant, the recommended systems should also be used for studying the basic components of the extraction processes, e.g. formation and motion of drops, transfer of heat and mass, coalescence, interphase phenomena, etc. Hopefully, the work of different authors could then become comparable and the explanation of anomalous phenomena, deviations and strange behaviour will be possible.

The Sub-Committee of the Working Party faced a difficult task in selecting the systems which should be considered and eventually recommended for general use after thorough investigation. The selection necessarily involved compromises and the systems chosen for this study are not ideal. However, they can undoubtedly meet the objective and are supported by the experience of the principal institutions dealing with liquid extraction in Europe, the data in the literature and the results of the contributors obtained during this project. The following systems were chosen for thorough investigation:

toluene - water - acetone
n-butanol - water - succinic acid
methyl isobutyl ketone - water - acetic acid.

For practical reasons they are all aqueous but they do differ in the most significant physical property for two-phase liquid systems - interfacial tension. Systems (1) and (2) are characterized by distribution coefficients approaching unity and constant over a wide concentration range. They may be assumed to involve physical extraction in contrast to System (3) which, while being widely used in practice, show some anomalous behaviour. System (1) represents an essentially immiscible basic binary system, whereas in Systems (2) and (3) the basic binaries are mutually soluble. The characters of Systems (1) and (3) provide very different wettability of different solids by both phases, whereas in System (2) these differences are smaller. Analysis of all the systems chosen is simple; the results are reproducible and all the systems consist of easily available raw materials.

These arbitrarily chosen characteristics of the selected systems do not mean that it would be impossible to find other suitable or even more suitable systems. However, selection, description and introduction into practice would present difficulties which cannot easily be overcome at the present time.

This publication presents results of the work of a wide group of authors and is aimed at engineers as well as scientists dealing with extraction systems. Hence, an arrangement was accepted which makes possible utilisation of the data measured either for technical purposes in the form of a smoothed analytical function or in the original form for deeper studies. For the most part, the data have been measured by two or more contributors, using different methods and raw materials from different sources. The authors believe that this published set of data will encourage the search for a deeper interpretation on a physico-chemical basis.

The information on fire and safety properties and the behaviour of selected systems has been included for guidance only; the reader should check the appropriate national regulations, the observance of which is obligatory.

The text of the work has been kept as brief as possible. More detailed data on the measurements can be obtained from individual authors, whose names and addresses are presented.

October 1977

Ing. T. Mišek, DrSc.

Chairman of the Working Party Sub-Committee

List of Contributors

This publication originated from the joint work of many European experts. It would be invidious to differentiate between their contributions and so they are listed alphabetically:

L. Alders R. Berger H.W. Brandt J. Bulicka G.A. Davies J. Gmehling A. Hackl M. Hafez C. Hanson S. Hartland H. Hartmann J. Kos J. v.d. Leur A. Newton U. Onken V. Pavasovic J. Prochazka V. Rod H. Sawistowski J. Schröter M. Soika W. Solar G. Strauss W.D.B. von Wolfersdorff

G. Ziebland

l

The Netherlands Federal Republic of Germany Federal Republic of Germany Czechoslovakia United Kingdom Federal Republic of Germany Austria Switzerland United Kingdom Switzerland Federal Republic of Germany The Netherlands The Netherlands United Kingdom Federal Republic of Germany Yugoslavia Czechoslovakia Czechoslovakia United Kingdom Federal Republic of Germany Federal Republic of Germany Austria Federal Republic of Germany Federal Republic of Germany Austria