

An Engineering Geological Classification of Limestone Material

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P.D. Swart



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<p>This report is a thesis study submitted to Professor D.G. Price of the Section Engineering Geology of the Department Mining and Petroleum Engineering at the Delft University of Technology in conformity with the requirements for the degree of Mijningenieur (Mining Engineer).</p>	
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AN ENGINEERING GEOLOGICAL CLASSIFICATION
OF LIMESTONE MATERIAL

BY P. D. SWART

November 1937

This report is a thesis study submitted to Professor B. D. Price of the Section Engineering Geology of the Department Mining and Petroleum Engineering at the Gulf University of Technology in conformity with the requirements for the degree of M. S. (Mining Engineer).

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Abstract

In engineering practice there exists no complete classification for pure limestone material. Although there are a number of classification systems, which rely on useful parameters, the problem is that there are no indications of quality regarding the requirements of the various engineering disciplines. In order to develop a classification system and to obtain a better insight in the behaviour of limestone, samples of twelve types of limestone from different geological formations were extensively tested with both mechanical and physical techniques. A simple test to indicate the solubility of a limestone can be performed with hydrochloric acid. Based on the results of the tests an empirical classification system is proposed for four engineering fields who are concerned with limestone: Tunelling, Foundation, Excavation and Aggregates.

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Abstract

In engineering practice there exists no complete classification for pure limestone material. Although there are a number of classification systems, which rely on useful parameters, the problem is that there are no definitions of quality regarding the relationships of the various engineering disciplines. In order to develop a classification system and to obtain a better insight in the behavior of limestone, samples of twelve types of limestone from different geological formations were extensively tested with both mechanical and physical techniques. A simple test to indicate the solubility of limestone can be performed with hydrochloric acid. Based on the results of the tests an empirical classification system is proposed for four engineering fields who are concerned with limestone: Tunneling, Foundation, Excavation and Aggregates.

1. Introduction

Upon searching the literature for a classification of carbonate rocks it was discovered that none of the existing classifications of carbonate rock material seem to render any satisfactory information as to the engineering behaviour of these rocks.

The existing classifications appear to be developed only for a single purpose, they were required to serve. Generally the complexity and confusion of these classifications contribute to a distortion of their engineering quality. Consequently the parameters of such a system, which can be of geological or engineering origin, can not render enough clear information for complete understanding of the behaviour of the rocks.

Acknowledgements

The realisation of this study has been a task which could not have been done without valuable support to solve the many problems encountered.

Especially I wish to thank Prof. D.G. Price, who has given inexpressible advice regarding the achievement of the classification system. Secondly Ing. W. Verwaal has my special gratitude for his technical support during the mechanical testing and for the use of his excellent data acquisition programs.

Furthermore I thank A.A. van Dijk for his help with the preparation of the samples and the thin sections; W. de Grauw and Th. Verkroost for their help with the determination of the calcium- and magnesium-carbonate contents; R. de Haas for his help with the determination of the porosity; the quarry managers, who gave permission to sample the limestone blocks; and the members of the staff of the Engineering Geology section, who helped and advised me.

1.1 Geological Background

For full comprehension of the classification systems for carbonate rocks it is necessary to understand the factors that influence the origin and formation of these rocks.

Pure limestones, which are generally considered to be those carbonates composed of more than 99% calcium carbonate CaCO_3 (Fig. 2.2.1. Peckes & Higginbottom 1978), are a group of sedimentary rocks, which occur in a wide variety of forms and degrees of induration. The minerals calcite and aragonite (also CaCO_3), both for the main part chemical precipitates from seawater, form the basis of the major components of these rocks. Aragonite is an unstable combination, which transforms after consolidation into calcite. Aside from this the minerals dolomite

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1. Introduction

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The existing classifications appear to be developed only for a singular purpose, they were required to serve. Generally the complexity and confusing nomenclature of these classifications contribute to a limited interpretation of their engineering quality. Consequently the parameters of such a system, which can be of geological or engineering origin, can not render enough clear information for complete understanding of the behaviour of the carbonate material.

Apparently in all of these systems there seems to be no direct relationship between the amount of calcium carbonate (CaCO_3) present in the rock and its geotechnical properties (Datta, 1982). This hypothesis should make it rather difficult to distinguish limestone from other types of rock. The aim of this study is then to develop a classification system which will relate the different carbonate rocks in such a way that the information conveyed is of practical use to the engineering disciplines concerned.

In order to achieve this end the physical and mechanical properties of a number of European limestones were compared and contrasted. The properties determined have been chosen in relation to the requirements of the engineering disciplines involved with foundation, construction and excavation of the massive material of carbonate rocks.

1.1 Geological Backgrounds

For full comprehension of the classification systems for carbonate rocks it is necessary to understand the factors that influence the origin and formation of these rocks.

Pure limestones, which are generally considered to be those carbonates composed of more than 90% calcium carbonate CaCO_3 (fig. 2.2.1 Fookes & Higginbottom 1975), are a group of sedimentary rocks, which occur in a wide variety of forms and degrees of induration. The minerals calcite and aragonite (also CaCO_3), both for the main part chemical precipitates from seawater, form the basis of the major components of these rocks. Aragonite is an unstable combination, which transforms after consolidation into calcite. Aside from this the minerals dolomite

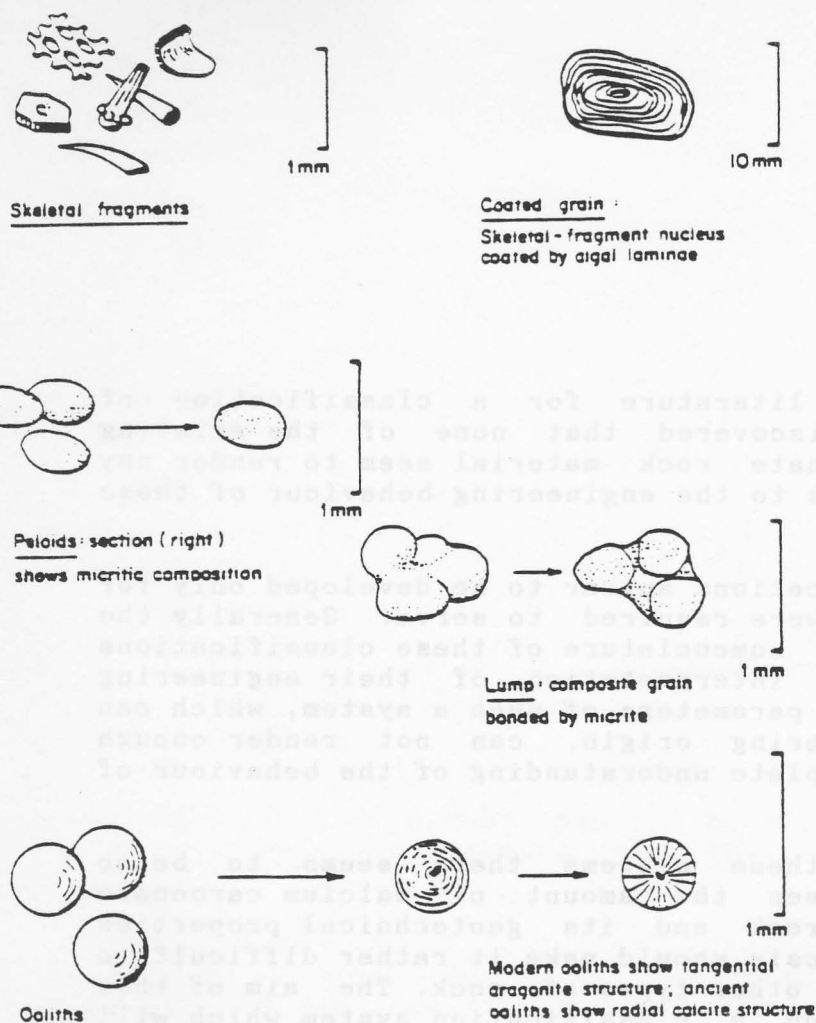
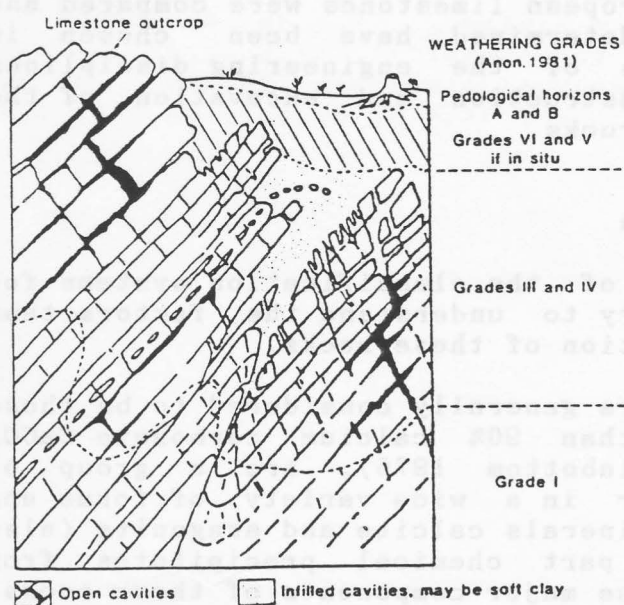


figure 1.1.1 Constituents of carbonate rocks



A typical weathering profile for carbonate rocks (modified from Deere and Patton, 1971,

figure 1.1.2 Weathering profile for carbonate rocks (Deere & Patton 1971)

	Great Limestone	Four Fathom Limestone
R.Q.D. %	79 - 95	82 - 97
Weathering	Unweathered - moderately	Unweathered
Rock Strength	100 - 200 MPa	100 - 200 MPa
Joint Spacing	0.3 - 1 m	1 - 3 m
Joint Separation	5 mm	0.1 - 1 mm
Joint Continuity	Continuous no gouge	Continuous with gouge
Number of localities	10	5
Rock Class	II - III Good, Fair	II Good
Stand-up Time	6 months - 1 week	6 months

Application of the CSIR (Bieniawski, 1973) classification to the limestones in the Kielder tunnels, N.E. England.

	Great Limestone	Four Fathom Limestone
R.Q.D. %	79 - 95	82 - 97
Jn	3 - 4	4 - 15
Jr	1 - 3	1,5 - 2,0
Ja	1 - 4	0,75- 1,0
S.R.F.	2,5 - 5,0	1 - 5,0
Rock quality	Q = 7 - 16,5	Q = 1,1 - 16,3

Application of the NGI (Barton et al., 1974) classification to the limestones in the Kielder tunnels, N.E. England. Jn is joint set number; Jr is joint roughness number; Ja is joint alteration number; SRF is stress reduction factor.

figure 1.2.1 Rock mass classification for limestone (1974,1973)

associated with pressure solution.

When consolidated carbonate rocks are exposed to the surface, they weather by solution, leaving behind the insoluble parts of the original rock. An extensive review of the mechanisms is given by Trudgill (1985). The rate of solution depends on the acidity of the water, the amount of CO_2 dissolved, the temperature, the waterflow, the previous saturation by CO_3 , and the solubility of the rock. When the surface area of the rock is exposed to such conditions weathering may occur in many forms. Deere and Patton (1971) have illustrated a typical weathering profile for carbonate rocks (fig 1.1.1). A description of weathering features of rocks recommended by the IAEG (the International Association of Engineering Geology (1970, 1972)) is presented in table 1.1.2 .

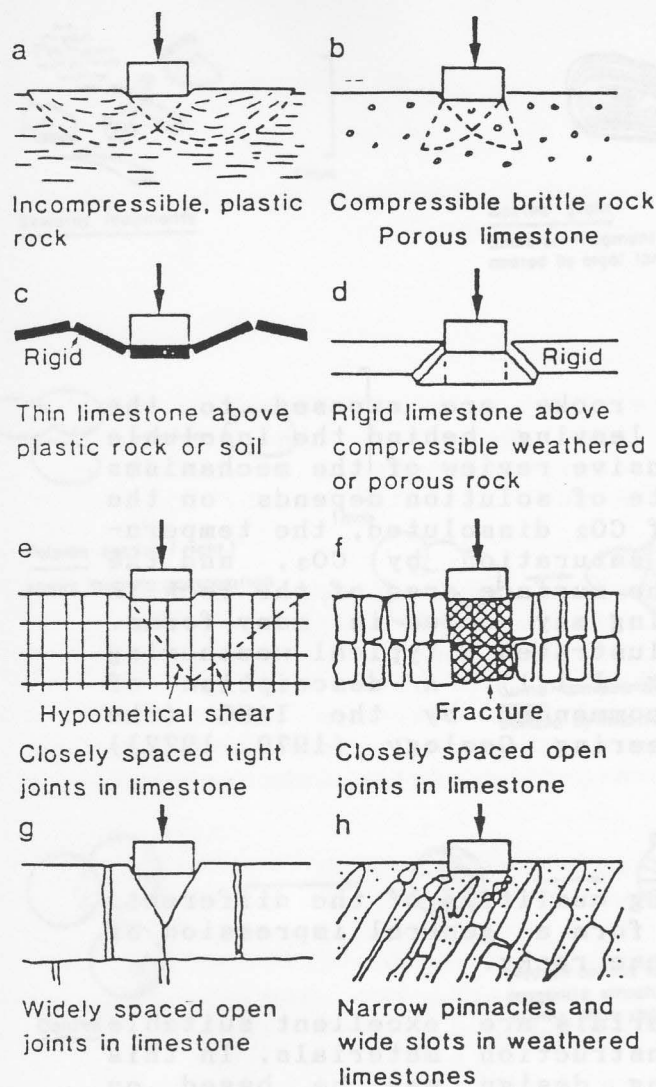
1.2 Engineering Considerations

An impression of the engineering qualities of the different carbonates may aid the reader to form a general impression of todays uses of and problems with these rocks.

The strong sound limestone materials are excellent suitable for foundations, tunnels and construction materials. In this category of rocks the engineering design may be based on classification systems for rock mass such as the CSIR (Bieniawski 1973) or the NGI (Barton 1974). An example applied on limestones is given in fig. 1.2.1 . A rock material classification is made by Deere and Miller (1966), who related the Unconfined Compressive Strength and the Elastic Modulus of intact rock (fig. 1.2.2).

Regarding the weakest carbonates, the spectrum ranges from slightly indurated soils to well cemented limestone or weathered and fractured limestone. The usefulness of cemented carbonate soils depends on the degree of induration and the percentage of carbonate material. The varying foundation conditions in this respect pose a serious problem. Sowers (1976) presented a qualitative approach to the types of foundation failures for limestones (figure 1.2.3). The degree of anisotropy, the influences of other media, and the discontinuity condition are accounted for. Hence the bearing capacity can be assessed from the compressibility of the limestone mass with the help of tests. Indications of the bearing capacity are noted in figure 1.2.4 according to the Code of Practice for Foundations (1972).

Regarding the use of carbonates as construction material, the suitability of weak cemented limestones as road aggregate is discussed by Netterberg (1982). As an example of characteristic limestone testing techniques used in road aggregate practice figure 1.2.5 presents data of the British Road Research Laboratory. The suitability of limestones as building stone requires a certain strength and is reversely proportional to the amount of fractures present in the rock. The physical and



Foundations on limestones. Varying modes of failure by:

- (a) General shear; (b) Local shear;
- (c) Slat action; (d) Punching shear;
- (e) Rankine shear; (f) Unconfirmed compression;
- (g) Splitting; (h) Local crushing.

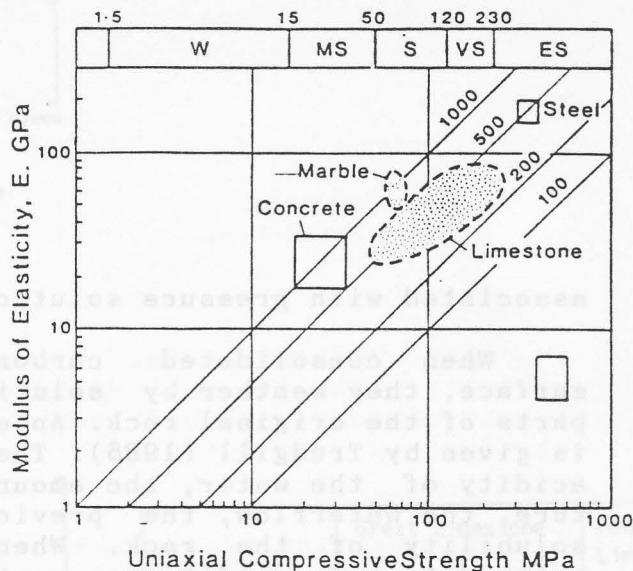
figure 1.2.3 Varying foundation conditions in limestones (Sowers 1976)

PHYSICAL PROPERTY	DENSITY OF LIMESTONE		
	Low	Medium	High
Absorption by weight, max, %	12	7.5	3
Density, min, kg/m ³	1760	2160	2560
Compressive strength, min, MPa	12	28	55
Modulus of rupture, min, MPa	2.9	3.4	6.9
Abrasion resistance, min, hardness*	10	10	10

*Relates to stone subject to foot traffic.

ASTM C568—79. Physical requirements for limestone building stone.

figure 1.2.6 requirements for limestone buildingstone (ASTM-C568 79)



A general engineering classification of limestones, dolomites and marbles in terms of elastic modulus and unconfined compressive strength (after Deere and Miller, 1966).

figure 1.2.2 A rock material classification for limestone (Deere & Miller 1966)

CLASS	ROCK TYPE	PRESUMED BEARING VALUE in kN/m ²	REMARKS
2	Hard limestones	4 000	These values are based on the assumption that the foundations are carried down to unweathered rock
6	Hard sound chalk, soft limestone	600	
7	Thinly bedded limestones	to be assessed after inspection	

Presumed bearing values for limestones under vertical static load from CP2004: 1972, Code of Practice for Foundations (Anon. 1972).

figure 1.2.4 Indication of bearing capacity (Code of Practice for foundations 1972)

and siderite often occur in these rocks as transformation products of calcium carbonate.

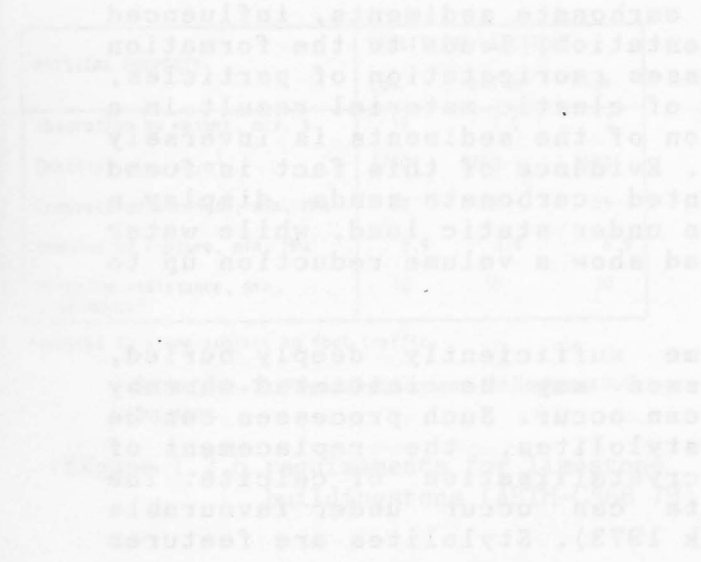
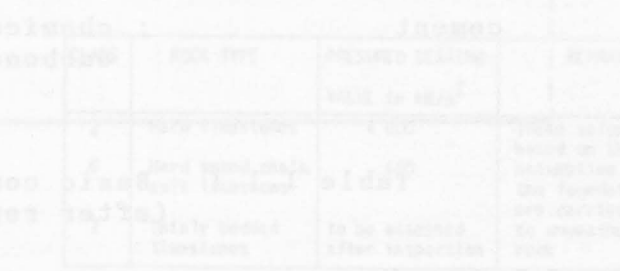
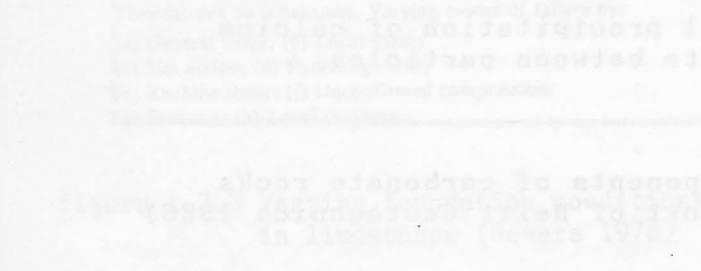
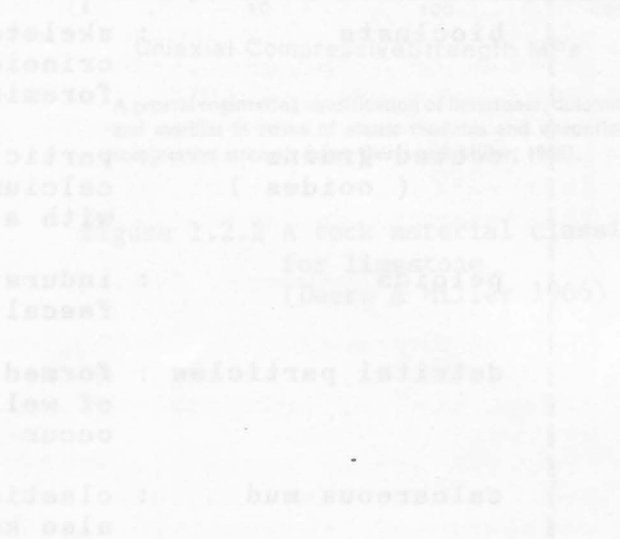
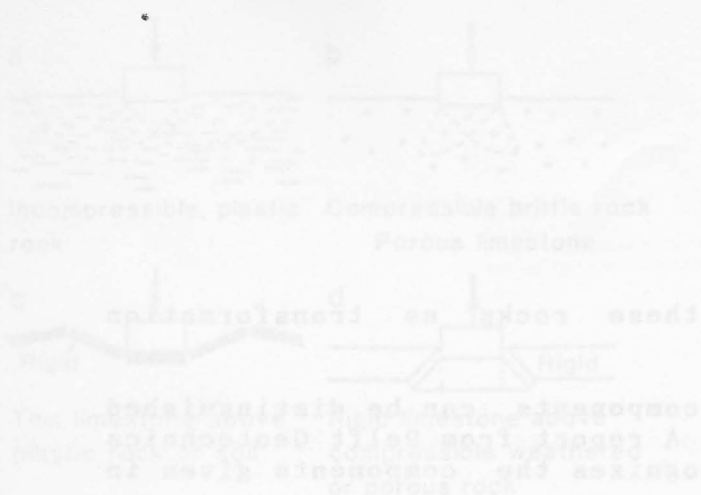
The origin of the carbonate components can be distinguished with reference to their fabric. A report from Delft Geotechnics (1986) on carbonate sediments recognizes the components given in table 1.1.1 .

bioclasts	: skeletal fragments formed by corals, crinoids, algae, brachiopods and foraminiferae etc.
coated grains (ooides)	: particles with a layered frame of calcium carbonate; when concentric and with a kernel they are called oolites
peloids	: indurated homogeneous spheres of faecal origin
detrital particles	: formed by mechanical disintegration of well consolidated carbonate rocks; occur in conglomerates and breccia
calcareous mud	: clastic products of micro organisms also known under the name of micrite
cement	: chemical precipitation of calcium carbonate between particles

Table 1.1.1 Basic components of carbonate rocks
(after report of Delft Geotechnics 1986)

Compaction and induration of carbonate sediments, influenced by temperature and previous cementation, leads to the formation of limestone. During these processes reorientation of particles, solution of lime and fracturing of clastic material result in a decreasing porosity. The compaction of the sediments is inversely proportional to their grainsizes. Evidence of this fact is found in arid climates where early cemented carbonate sands display a relative low degree of compaction under static load, while water saturated carbonate muds under load show a volume reduction up to 90 % .

Should the limestone become sufficiently deeply buried, diagenetic and metamorphic processes may be initiated whereby significant changes in the rock can occur. Such processes can be recognized by the presence of stylolites, the replacement of calcite by dolomite, and the recrystallisation of calcite. The replacement of calcite by dolomite can occur under favourable conditions of salinity (Pannekoek 1973). Stylolites are features



Should the limestone become sufficiently deeply buried, diagenetic and metamorphic processes may be superimposed on the original diagenetic changes in the rock. Such processes would be recognized by the presence of stylolites, the replacement of calcite by dolomite, and the replacement of calcite by dolomite. Under favorable conditions of salinity (Fennel 1973), stylolites are features

Sample No.	Crushing value	Aggregate impact value	Aggregate abrasion value	Water absorption (per cent)	Specific gravity	Polished stone coefficient
104	54 (17-21)	23 (17-32)	13.7 (7-22)	1.0 (0.5-2.5)	2.5 (2.3-2.8)	0.43 (0.30-0.73)
105	54	23	13.7	1.0	2.5	0.43

mechanical requirements for limestones as building stone are indicated in the ASTM C568-79 report (figure 1.2.6). Quality investigation could then focus on crushing characteristics, the degree of fissuring and the behaviour of minor fractions present in the material (Ashurst and Dimes 1977).

A major problem with the moderately weak to strong limestones is the solution effect, which results in the formation of cavities. Without adequate investigation of the limestone mass the risk of collapse is may be too high. From an engineering point of view the important factors to be assessed are the size and distribution of the fissures and the solution rate constant as defined by James (1982).

2 Existing Classifications of Carbonate Rocks

The factors that play a dominant role in the classification systems of carbonate rocks appear to depend largely on the purpose the classifications were designed to serve. A large number of classifications were developed because of the interest of the oil industry in the porosity and permeability of these rocks. Their aim was mainly to correlate the texture and depositional origin of the limestones to porosity values.

Upon reviewing the existing classifications it was found that they can be grouped into one of three categories ; geological, geotechnical or engineering geological classifications.

2.1 Geological Classifications

The two systems most commonly used for the classification of pure carbonates are those proposed by Dunham (1962) and Folk (1959, 1962).

The main criterium in Dunham's classification is the relation between particles and matrix. The terms packstone, grainstone, mudstone etc. should qualify the rock ranging from grain to matrix supported. Although the textural basis provides an insight in the depositional environments, it offers no room for an indication of the induration degree (fig 2.1.1).

Folk's classification is based on three elements ; particles (allochems), microcrystalline ooze (orthochems) and sparry (coarse) calcite cement. Each class may then be further divided giving an insight into their depositional maturity. By combining texture with the type of constituent a sequential nomenclature is developed (fig. 2.1.2). This leads to rock descriptions such as intra-bio-pel-micrite. This system is more suited to microscope studies than field use. As for the nomenclature according to grain size he presented a classification (fig. 2.1.3), which is based on the Wentworth scale. A weak point in his classification is that cementation degrees can not be derived.

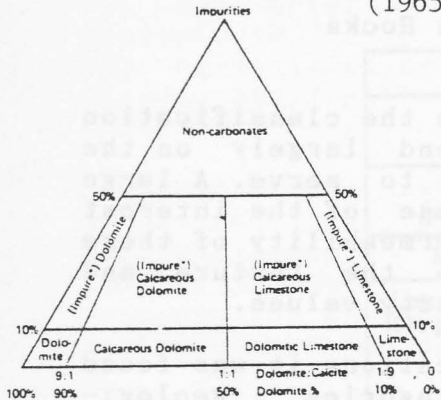
Schmidt (1965) tackled the spectrum between limestone and dolomite in extension of the division made by Petijohn (1959) (fig. 2.1.4).

Leighton and Pendexter (1962) proposed a classification considering three variables : grain size, the proportions of the matrix to the allochems (particles) and a terminology for the composition of carbonate rocks. The grainsizes of the different allochems are compared to the type of allochem, thus classifying the texture, while the geological descriptive terms, like an argillaceous limestone, supply information regarding the

Volume percent of rock	Detailed Classification	Standard Classification
Traces - 2 vol %	Extremely slightly dolomitic	Dolomitic
2 - 5 vol %	Very slightly dolomitic	
5 - 10 vol %	Slightly dolomitic	
10 - 25 vol %	Fairly dolomitic	
25 - 50 vol %	Highly dolomitic	

(b) Schmidt's detailed classification of calcite-dolomite mixtures.

Classification of limestone-dolomite mixtures figure 2.1.4 (Schmidt) (1965)



*An appropriate compositional term should be substituted for the word "impure"

Leighton and Pendexter's compositional terminology for carbonate rocks

figure 2.1.5 (Leighton & Pendexter 1962)

TABLE GRAIN-SIZE SCALE FOR CARBONATE ROCKS

		Transported Constituents	Authigenic Constituents		
64	mm	Very coarse calcirudite	Extremely coarsely crystalline	4	mm
16	mm	Coarse calcirudite			
4	mm	Medium calcirudite			
1	mm	Fine calcirudite	Very coarsely crystalline	1	mm
0.5	mm	Coarse calcarenite	Coarsely crystalline	0.25	mm
0.25	mm	Medium calcarenite			
0.125	mm	Fine calcarenite	Medium crystalline	0.062	mm
0.062	mm	Very fine calcarenite			
0.031	mm	Coarse calcilutite	Finely crystalline	0.016	mm
0.016	mm	Medium calcilutite			
0.008	mm	Fine calcilutite			
0.004	mm	Very fine calcilutite	Aphanocrystalline	0.004	mm

Carbonate rocks contain both physically transported particles (oolites, intraclasts, fossils, and pellets) and chemically precipitated minerals (either as pore-filling cement, primary ooze, or as products of recrystallization and replacement). Therefore, the size scale must be a double one, so that one can distinguish which constituent is being considered (e.g., coarse calcirudites may be cemented with very finely crystalline dolomite, and fine calcarenites may be cemented with coarsely crystalline calcite). The size scale for transported constituents uses the terms of Grabau but retains the finer divisions of Wentworth except in the calcirudite range; for dolomites of obviously allochemical origin, the terms "dolorudite," "dolarenite," and "dolojutite" are substituted for those shown. The most common crystal size for dolomite appears to be between .062 and .25 mm and for this reason that interval was chosen as the "medium crystalline" class.

figure 2.1.3 (Folk 1962)

LEIGHTON & PENDEXTER'S TEXTURAL CLASSIFICATION OF LIMESTONES

Grain/micrite ratio	% Grains	Grain type					Organic frame builders	No organic frame builders
		Detrital	Skeletal	Pellets	Lumps	Coated Grains		
9:1	90	Detrital Lst	Skeletal Lst	Pellet-Lst	Lump-Lst	Oolitic Lst Pisolithic Lst Algal encr Lst	Coralline Lst Algal Lst etc	
1:1	50	Detrital Micritic Lst	Skeletal Micritic Lst	Pellet-Micritic Lst	Lump-Micritic Lst	Oolitic (Pisolithic etc) Micritic Lst	Coralline-micritic Lst Algal-micritic Lst etc	
1:9	10	Micritic-Detrital Lst	Micritic-Skeletal Lst	Micritic-Pellet Lst	Micritic-Lump Lst	Micritic-oolitic (Pisolithic etc) Lst	Micritic-coralline, Micritic-algal Lst etc	
Micritic Limestone								

Figure 2.1.2 ab a simplified scheme (Folk 1962)

Percent Particles	OVER 2/3 LIME MUD MATRIX (Micrite)				SUBSEQUENT SPAR AND LIME MUD	OVER 2/3 SPARRY CEMENT (Sparrite)		
	0-1%	1-10%	10-50%	> 50%		SORTING POOR	SORTING GOOD	ROUNDED & ABRASSED
Representative Rock Terms	MICRITE & DISMICRITE	FOSSILIFEROUS MICRITE	SPARSE BIOMICRITE	PACKED BIOMICRITE	POORLY WASHED BIOSPARITE	UNSORTED BIOSPARITE	SORTED BIOSPARITE	ROUNDED BIOSPARITE
Terminology Folk 1959	Micrite & Dismicrite	Fossiliferous Micrite	Biomicrite		Biosparite			
Terrigenous Analogues	Claystone		Sandy Claystone	Clayey or Immature Sandstone	Submarine Sandstone	Mature Sandstone	Supermature Sandstone	

■ LIME MUD MATRIX

▨ SPARRY CALCITE CEMENT

percentages of limestone, dolomite and impurities (fig. 2.1.5).

A Classification in which the porosity plays a significant role is represented in the unified system of Choquettte & Pray (1970). In this system an important distinction is made between the terms "fabric selective" and "not fabric selective" porosity. The term fabric selective means that pores are bounded by particle boundaries. The system is designed for carbonate rocks but is only a partial description (fig. 2.1.6).

2.2 Geotechnical Classifications



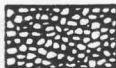




Due to rapid expansion of engineering developments during the 70's, one of the first geotechnical oriented classifications was developed by Fookes and Higginbottom (1975). Their classification of near-shore carbonate rocks included a degree of induration, grain size, mineral composition and the origin of the carbonate material. They connected differences in strength and grain size to common known names for pure and impure limestones. This classification formed a basis for many modifications and extensions (fig. 2.2.1 a). Apart from the classification for engineering purposes, they defined a nomenclature for pure and impure carbonate sediments (fig. 2.2.1 b & c).

Clark and Walker (1977) modified the nomenclature and encompassed the impure carbonates as well, still using the same four variables as Fookes and Higginbottom. The engineering behaviour was quantified by boundaries using ranges of approximate Unconfined Compressive Strength (UCS) values in relation to the degree of induration (fig. 2.2.2).

For slightly cemented sands Beringen (1982) adjusted the Clark & Walker system to the use of the cone penetration test. The link between cone resistance and the cementation is quantitative described for indurated carbonate soils (fig. 2.2.3).

Regarding pure carbonate sediments the Clark and Walker classification was modified by Fugro (1978) for the North Rankin Project in Australia (fig. 2.2.4). Mainly concerned with cemented soils this classification discerns cementation and induration; the induration should be the result of cementation and cohesion. The parameters used are grainsize, name, degree of cementation, bedding and lamination, origin of carbonate, colour, and the minor fractions.

With regard to carbonate sands in India, Datta (1982) modified the Clark & Walker classification and stated some remarks for a new classification. Although Datta is not directly involved with limestones, the cemented carbonate sands he is concerned with overlap the spectrum of weak to very weak limestones. According to his paper, aspects which could supply

BASIC POROSITY TYPES			
FABRIC SELECTIVE		NOT FABRIC SELECTIVE	
	INTERPARTICLE	BP	
	INTRAPARTICLE	WP	
	INTERCRYSTAL	BC	
	MOLDIC	MO	
	FENESTRAL	FE	
	SHELTER	SN	
	GROWTH-FRAMEWORK	GF	

MODIFYING TERMS			
GENETIC MODIFIERS		SIZE* MODIFIERS	
PROCESS	DIRECTION OR STAGE	CLASSES	
SOLUTION s	ENLARGED e	MEGAPORE mg	large 256
CEMENTATION c	REDUCED r		small 32
INTERNAL SEDIMENT i	FILLED f	MESOPORE ms	large 4
			small 1/2
		MICROPORE mc	small 1/16
TIME OF FORMATION		Use size prefixes with basic porosity types: meSOL mg, meVUG msVUG, meMO msMO, meSN msSN, meGF msGF *For regular-shaped pores smaller than cavern size. †Measures refer to average pore diameter of a single pore or the range in size of a pore assemblage. For tubular pores use average cross-section. For platy pores use width and note shape.	
PRIMARY P		ABUNDANCE MODIFIERS	
pre-depositional Pp			
depositional Pd			
SECONDARY S			
egogenic Se		percent porosity	(15%)
mesogenic Sm		or	
telogenic St		ratio of porosity types	(1:2)
		or	
		ratio and percent	(1:2) (15%)
Genetic modifiers are combined as follows:			
PROCESS + DIRECTION + TIME			
EXAMPLES:			
solution-enlarged	se		
cement-reduced primary	crP		
sediment-filled egogenic	fSe		

figure 2.1.6 : Classification of the porosity of carbonate rocks (Choquette & Pray 1970)

useful information for the evolvement of a further classification system, are :

- cementation
- grainsize distribution and plasticity
- nature of carbonate components
- nature of non-carbonate components

To specify the geotechnical properties he suggested the following descriptive terms should be included in a classification system for carbonate soils :

- susceptibility to crushing
- degree of uniformity and cementation
- influence of carbonate material in relation to non-carbonate material

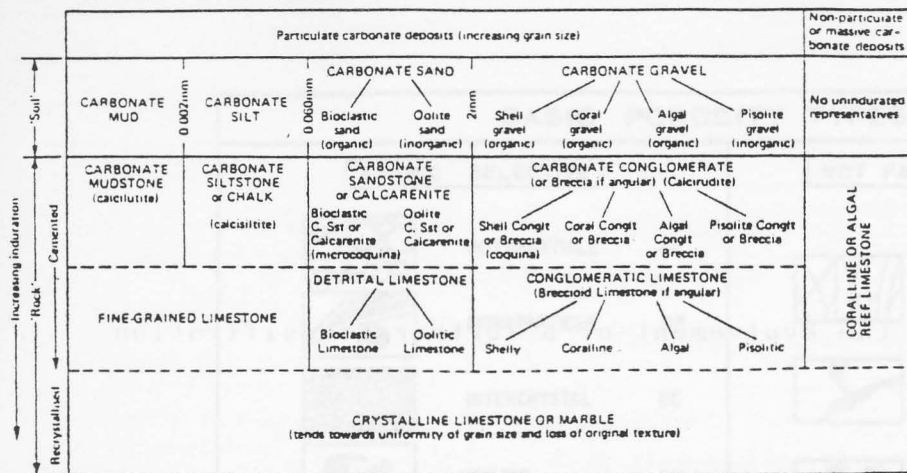
The scheme proposed by Datta for geotechnical description is listed in fig. 2.2.5 .

2.3 Engineering Geological Classifications

An engineering geological classification of carbonate rocks with a descriptive character was published by Burnett and Epps (1979). They suggested a double ternary diagram of pure limestone and admixtures in addition to individual classification tables for grain size, joint spacing, density, weathering, texture and strength. Unfortunately the terminology they propose becomes somewhat extensive. A description sheet which, could be used for the sampling of carbonate rocks, is given in appendix A.1 .

Dearman (1981) considered the engineering geological description system in which he accounts for carbonate rocks as a material as well as their rock mass. In his paper he recognizes the importance of different properties to the various engineering fields involved with limestone. Furthermore he shows that correlations exist between engineering index properties and geological properties. As example he presents a correlation between the engineering geological grade and the mechanical properties of chalk at Mundford (after Ward et al. 1968) in figure 2.3.1 This figure has been widely used as a preliminary estimate in the logging of drill cores and site investigation pits and trenches. Visual description then enables to derive deformation characteristics of the chalk, based on comparison with a known deformation and description.

FOOKES & HIGGINBOTTOM'S ENGINEERING CLASSIFICATION OF CARBONATES (established alternative names are in brackets)



Siltstone	Calcareous Siltstone	Silty Limestone	Limestone
10%	50%	10%	
Conglomerate	Calcareous Conglomerate	Conglomerate Limestone	Limestone

(a) Classification of indurated silt-carbonate and gravel-carbonate sediments.

0	Pure Limestone
5	
20	Marly Limestone
35	Limey Maristone
65	MARLSTONE
80	Clayey Maristone
95	Marly Claystone
100	Pure Claystone

(b) Classification of indurated clay-carbonate sediments.

Higginbottom and Fookes' compositional classification of impure carbonate sediments

- Non-carbonate constituents are likely to be siliceous apart from local concentrations of minerals such as feldspar and mixed heavy minerals (Emery 1956).
- In description the rough proportions of carbonate and non-carbonate constituents should be quoted and details of both the particle minerals and matrix minerals should be included.
- The preferred lithological nomenclature has been shown in block capitals; alternatives have been given in brackets and these may be substituted in description if the need arises.
- Calcareous is suggested as a general term to indicate the presence of unidentified carbonate. Where applicable, when mineral identification is possible calcareous referring to calcite or alternative adjectives such as dolomitic, aragonitic, sideritic etc. should be used.

DEGREE OF INDURATION	APPROXIMATE UNCONFINED COMPRESSIVE STRENGTH	ADDITIONAL DESCRIPTIVE TERMS BASED ON ORIGIN OF CONSTITUENT PARTICLES							
		NOT DISCERNIBLE	BIOLASTIC (Inorganic)	DOLITE (Inorganic)	SHELL (Organic)	CORAL (Organic)	ALGAL (Organic)	PSOLITES (Inorganic)	
INCREASING GRAIN SIZE OF PARTICULATE DEPOSITS									
		0-002mm	0-060mm	2mm	60mm				
Non-indurated	Very soft to hard (<36 to $>300\text{MN/m}^2$)	CARBONATE MUD	CARBONATE SILT	CARBONATE SAND	CARBONATE GRAVEL				90%
		Clayey CARBONATE MUD	Siliceous CARBONATE SILT ①	Siliceous CARBONATE SAND ①	Mixed carbonate and non-carbonate GRAVEL ②				50%
Slightly indurated	Hard to moderately weak (0.3 to 12.5MN/m^2)	CLAY	SILT	silica SAND	GRAVEL				10%
		CALCILUTITE (carb. clayst.)	CALCISILTITE (carb. silst.)	CALCARENITE (carb. sandst.)	CALCIRUDITE (carb. conglom. or breccia)				90%
Moderately indurated	Moderately strong to strong (12.5 to 100MN/m^2)	Clayey CALCILUTITE	Siliceous CALCISILTITE	Siliceous CALCARENITE	Conglomeratic CALCIRUDITE ②				50%
		Calcareous CLAYSTONE	Calcareous SILTSTONE	Calcareous SANDSTONE	Calcareous CONGLOMERATE				10%
Highly indurated	Strong to extremely strong (170 to $>200\text{MN/m}^2$)	CLAYSTONE	SILTSTONE	SANDSTONE	CONGLOMERATE OR BRECCIA				10%
		Fine-grained Argillaceous LIMESTONE	Fine-grained Siliceous LIMESTONE	Siliceous detrital LIMESTONE	Conglomeratic LIMESTONE ②				50%
		CLAYSTONE	SILTSTONE	SANDSTONE	CONGLOMERATE OR BRECCIA				10%
CRYSTALLINE LIMESTONE OR MARBLE (tends towards uniformity of grain size and loss of original texture)									50%
Conventional metamorphic nomenclature applies in this section									

Proposed classification chart for description of Middle Eastern sedimentary rocks

figure 2.2.2 : Classification of Middle Eastern sedimentary rocks (Clark & Walker 1977)

2.4 Restrictions on Classification for Engineering Purposes

A striking fact is the differentiation of classification systems with reference to the nature of the carbonate rocks or soils, to which the system was applied. For example the Clark & Walker system encompasses the Middle Eastern sedimentary rocks, while Datta used his description for slightly cemented soils in India. The problem is that the definition of what is rock and what is soil has not been unambiguously assessed for the carbonate sediments. Hence due to subjective interpretation of the boundaries, the nomenclatures of two such systems can cause confusion when they are applied to the same carbonate.

In pursuit of prediction of engineering behavior many different factors play roles of alternating importance, like ranges of induration where cemented soils must be treated differently than sound rocks. Therefore it is desirable that standard values should discriminate between important boundaries.

Effective prediction of engineering properties to various carbonate rocks can be enhanced by relating their engineering uses. According to the standard procedures the most required properties of the following engineering fields were compared.

- Underground Excavation (Bieniawski 1979 and OPAC 1987)
- Dredging (PIANC 1984)
- Foundation Engineering (Fugro 1978, Dearman 1981)
- Construction Materials (ASTM-C 33 and ASTM C 568 1979)

Strength, particle size and bulk density appear to be essential engineering properties and additional information may be supplied about crushing and deformation characteristics.

For strength several classifications exist, some with rather poor boundary restrictions. In general the most suitable seems to be the strength of intact rock developed by the Geological Society (1970) given in table 2.4.1.

very weak	0 - 1.25	MPa
weak	1.25 - 5	MPa
moderately weak	5 - 12.5	MPa
moderately strong	12.5 - 50	MPa
strong	50 - 100	MPa
very strong	100 - 200	MPa
extremely strong	>200	MPa

Table 2.4.1 Strength classification of the Geological Society (1970)

		FINE GRAINED					MEDIUM-COARSE GRAINED											
		INCREASING GRAINSIZE																
		0.002 mm	0.006 mm	0.02 mm	0.06 mm	0.2 mm	0.6 mm	2 mm	6 mm	20 mm	60 mm							
SOIL	DEGREE OF INDURATION	APPROX. UNCONF. COMPR. STRENGTH															DEGREE OF CEMENTATION	APPROX. UNCONF. COMPR. STRENGTH
	very soft	0 - 12.5 KN/m ²															uncemented	0 - 25 KN/m ²
	soft	12.5 - 25 KN/m ²															very weakly cemented	25 - 50 KN/m ²
	firm	25 - 50 KN/m ²															weakly cemented	50 - 100 KN/m ²
	stiff	50 - 100 KN/m ²															firmly cemented	100 - 300 KN/m ²
	very stiff	100 - 200 KN/m ²															well cemented	0.3 - 12.5 MN/m ²
ROCK	hard	200 - 300 KN/m ²															hard cemented	12.5 - 100 MN/m ²
	weak - mod. weak	0.3 - 12.5 MN/m ²															mod. strong - strong	12.5 - 100 MN/m ²
	strong - extr. strong	70 - >200 MN/m ²															strong - extr. strong	70 - >200 MN/m ²

INCREASING GRAINSIZE														
0.002 mm	0.006 mm	0.02 mm	0.06 mm	0.2 mm	0.6 mm	2 mm	6 mm	20 mm	60 mm					
fine	med	coarse	fine	medium	coarse	fine	medium	coarse						
CARBONATE MUD			CARBONATE SILT		CARBONATE SAND			CARBONATE GRAVEL						
					clastic/bioclastic/oolitic			clastic/bioclastic						
CALCILUTITE (carb. mudstone)			CALCISILTITE (carb. siltstone)		CALCARENITE (carb. sandstone)			CALCIRUDITE (carb. congl. or breccia)						
					clastic/bioclastic/oolitic			clastic/bioclastic						
FINE GRAINED LIMESTONE					DETRITAL LIMESTONE			CONGLOMERATE LIMESTONE						
					clastic/bioclastic/oolitic			clastic/bioclastic						
CRYSTALLINE LIMESTONE					CRYSTALLINE LIMESTONE									

CLASSIFICATION OF CARBONATE SEDIMENTS (90-100% carbonate)
(modified after Clark and Walker, 1977)

figure 2.2.4 (fugro 1987) geotechnical classification of carbonate sediments

DEGREE OF INDURATION CEMENTATION	CONE RESISTANCE q_c MN/m ²
VERY WEAKLY INDURATED CEMENTED	0 - 2
WEAKLY INDURATED CEMENTED	2 - 4
FIRMLY INDURATED CEMENTED	4 - 10
WELL INDURATED CEMENTED	> 10

figure 2.2.3 (Beringen 1982)
the relation between cone resistance and degree of cementation or induration.

TABLE Proposed system of description.

Description of	Remarks
1. Cementation (a) No cementation (b) Weak cementation (c) Strong cementation (i) uniform (ii) partial	the soil has a soft rock-like appearance. Unconfined compressive strength should be indicated the soil contains cemented aggregates—this should be noted
2. Grain Size Distribution (GSD) and Plasticity (a) Grain size distribution (b) Plasticity	for strongly cemented soils, GSD is not very relevant; for uniform cementation, size of constituent particles should be indicated; for partial cementation, GSD of soil after removing aggregates should be indicated and size and proportion of aggregates noted separately for fine-grained soils in which intraparticle voids cause error in GSD and Atterberg limits, field classification procedures may be used for providing the relevant information in a qualitative sense
3. Nature of Carbonate Component (a) Carbonate content (b) Particle size of carbonate material (c) Particle characteristics and origin (d) Mineralogy (e) Geologic name	soils having more than 30% carbonate content should be termed as carbonate soils the carbonate content in the sand and in the silt-clay fractions should be determined separately and indicated. Microscopic studies mentioned below will also give information about particle size microscopic studies—optical microscope for sands and scanning electron microscope for fine-grained soils—should be conducted. Presence of thin-walled material and intraparticle voids should be highlighted X-ray diffraction analysis should be performed if possible to identify, the geologic name may be indicated
4. Nature of Noncarbonate Component (a) Particle size (b) Particle characteristics (c) Mineralogy	information on noncarbonate material is determined by dissolving the carbonate material in HCl, separating the remaining soil, and conducting the following tests on it grain size distribution analysis microscopic studies X-ray diffraction analysis

figure 2.2.5 Proposed system of description (Datta 1982) of carbonate soils

For dominant particle sizes the norms of MIT, DIN 4023, BS 1377 and the PIANC agree at convenient boundaries. Even more they are the same boundaries used by Clark & Walker and are therefore assumed to be suitable to carbonate rocks (table 2.4.2).

cobbles	200	-	60	mm
gravel	60	-	2	mm
sand	2	-	0.06	mm
silt	0.06	-	0.002	mm
clay		<	0.002	mm

Table 2.4.2 Particle Size boundaries used by BS 1377, DIN 4023, MIT and the PIANC.

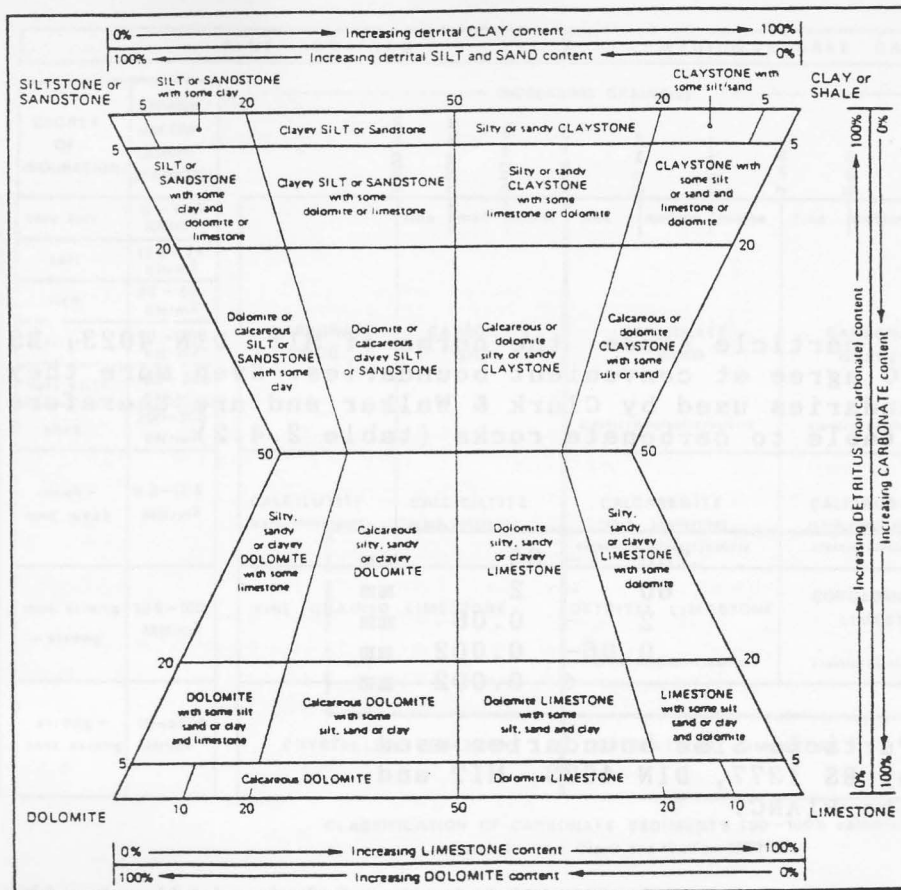
As in soil classifications it is useful to indicate the grainsize distribution. In most cases for limestones a distinction between particle and matrix sizes and percentages is probably generally sufficient.

Concerning bulk density the existing ranges are indicated for limestone building stone by the ASTM (American Society of Testing Materials) as given in table 2.4.3. The quality of a building stone is not directly related to the density.

Low density	:	1.76 - 2.16 (g/cm ³)
Medium density	:	2.16 - 2.56 (g/cm ³)
High density	:	2.56 - (g/cm ³)

table 2.4.3 Density division for limestone building stone (ASTM C568-79)

Complementary to the engineering parameters the important geological factors which were recognized are texture and mineral composition. Both these factors give a qualitative indication for parameters such as abrasion, drillability and cuttability. The description of carbonate and non-carbonate constituents is of great significance to the engineering behaviour of the rock. For example in limestones the presence of quartz does increase their abrasive capacity and clay minerals may cause swelling. Textural division must distinguish between clastic particles and crystal-line components.



Proposed classification of pure and admixtures of carbonate rocks

figures 2.3.1 a. t/m f (Burnett & Epps 1979)

STRENGTH CLASSIFICATION

engineering geological classification of carbonate rocks

Field Definition	Extrudes between fingers when squeezed	Very easily moulded with fingers	Moderate finger pressure required to mould	Moulded only by strong finger pressure	Cannot be moulded with fingers	Brittle or very tough	Crumbles in hand	Thin slabs break easily in hand	Thin slabs broken by heavy hand pressure	Lumps or core broken by light hammer blows	Lumps or core broken by heavy hammer blows	Lumps only chip by heavy hammer blows. Dull ringing sound	Rocks ring on hammer blow. Sparks fly
Description	Very soft	Soft	Firm	Stiff	Very Stiff	Hard	Very Weak	Weak	Moderately Weak	Moderately Strong	Strong	Very Strong	Extremely Strong
Strength Categories	Shear Strengths of Clays (kN/m ²) +						Unconfined compressive Strengths of Rocks (MN/m ²)						
	18	36	72	144	288		1.25	5	12.5	50	100	200	
							Point Load Strengths of Rocks (MN/m ²)						
							0.075	0.3	0.75	3	6	12	

* Based on the approximate relation:- Comp. Strength = 16 Point Load Comp. Strength
+ Various published strength categories exist — quote reference used

SIZE, SPACING, DENSITY AND PLASTICITY CLASSIFICATIONS

Soil & Rock Discontinuity Spacing Descriptions											Extremely Narrow	Very Narrow	Narrow	Mod. Wide	Wide	Very Wide	
Bedding Spacing Descriptions											Thinly Laminated	Thickly Laminated	Very Thin	Thin	Medium	Thick	Very Thick
Rock Grain Size Descriptions	Very Fine	Fine			Medium			Coarse			Very Coarse						
Soil Particle Size Divisions	Clay	Silt			Sand			Gravel			Cobbles	Boulders					
		Fine	Medium	Coarse	Fine	Medium	Coarse	Fine	Medium	Coarse							
mm 0.002 0.006 0.02 0.06 0.2 0.6 2 6 20 60 200 600 2000																	
Diameter or Spacing																	
Relative Density of Granular Soils		%	Very Loose		15	Loose		35	Medium Dense		65	Dense		85	Very Dense		Relative Density N Values
		Blows			4			10			30			50			
Plasticity	Field Estimate	Lean			Intermediate			Fat			Very Fat			Extra Fat			
Descriptions	Laboratory Term	Low			Intermediate			High			Very High			Extra High			
Liquid Limit, %		20		35		50		70		90 Liquid Limit, %							

PROPOSED TEXTURAL CLASSIFICATION OF CARBONATES

Depositional texture recognisable					Depositional texture not recognisable
Containing Micrite matrix			Grain supported – lacking micrite matrix	Original components bound together	
Matrix supported		Grain supported			Crystalline Limestone
Less than 10% grains	More than 10% grains	Oolitic, Skeletal, Shelly, or Detrital – micritic Limestone	Algal, Coralline, etc. Limestone		
Micritic Limestone	Micritic – Oolitic, Skeletal, Shelly, or Detrital Limestone				

figures 2.3.1 d t/m f
engineering geological classification
of carbonate rocks
(Burnett & Epps 1979)

WEATHERING CLASSIFICATION

Soil Mass			Rock Mass		
Diagnostic Features	Term	Grade Symbol	Grade Symbol	Term	Diagnostic Features
			W VI	Residual Soil	Rock is discoloured and completely changed to a soil in which original rock fabric is completely destroyed.
Soil discoloured and totally altered, with no trace of original structures.	Completely Weathered	W V	W V	Completely Weathered	Rock is discoloured and changed to a soil but original fabric is mainly preserved. Occasional small corestones.
Soil mainly altered with occasional small lithorelics of original soil. Little or no trace of original structures.	Highly Weathered	W IV	W IV	Highly Weathered	Rock is discoloured; discontinuities may be open and surfaces discoloured. Original rock fabric near discontinuities may be altered and penetrate deeply inwards but corestones are still present.
Soil composed of large discoloured lithorelics or original soil separated by altered material. Alteration penetrates inwards from discontinuities.	Moderately Weathered	W III	W III	Moderately Weathered	Rock is discoloured; discontinuities may be open and will have discoloured surfaces with alteration starting to penetrate inwards. Intact rock noticeably weaker than fresh rock.
Material composed of angular blocks of fresh soil which may or may not be discoloured. Some alteration starting to penetrate inwards from discontinuities separating blocks.	Slightly Weathered	W II	W II	Slightly Weathered	Rock may be slightly discoloured, particularly adjacent to discontinuities, which may be open and will have slightly discoloured surfaces. Rock not noticeably weaker than fresh rock.
			W IA	Faintly Weathered	Weathering limited to the surfaces of major discontinuities.
Parent soil shows no discolouration or loss of strength. Discontinuities usually tight and not discoloured.	Fresh	W I	W IA	Fresh	Parent rock shows no discolouration or loss of strength. Discontinuities usually tight and not discoloured.

DIAGNOSIS OF CARBONATE TYPES WITH DILUTE HYDROCHLORIC ACID

(Test comprises immersing rock chips about $\frac{1}{2}$ in dia \times $\frac{1}{2}$ in thick in cold dilute acid)

Rock Type	Reaction with cold dilute HCL (1:7:10)
Limestone	Violent effervescence; frothy audible reaction; small chips will bob about and tend to float on the surface
Dolomitic Limestone	Brisk, quiet effervescence; small chips skid about on the bottom of the container and rise slightly off the bottom; there is a continuous stream of CO ₂ through the acid.
Calcitic Dolomite	Mild emission of CO ₂ ; beads; small specimens may vibrate, but tend to stay in one place.
Dolomite	No effervescence, no immediate reaction; slow formation of CO ₂ ; beads on the surface of the rock; reaction slowly accelerates until a thin stream of beads rises to the surface, especially when heated.

Grade	Brief description	Approximate range of E (kg/sq.cm)	Bearing pressure Causing 'yield' (kg/sq.cm)	Creep properties
V	Structureless remoulded chalk containing lumps of intact chalk	<5000	<2	Exhibits significant creep
IV	Friable to rubbly chalk with open joints often infilled with soft remoulded chalk	5000-10 000	2-4	Exhibits significant creep
III	Medium to hard rubbly to blocky chalk with closely spaced slightly open joints	10 000-20 000	4-6	For pressures not exceeding 4 kg/sq. cm creep is small and terminates in a few months
II	Medium hard to hard chalk with widely-spaced, tight joints	20 000-50 000	10	Negligible creep for pressures of at least 4 kg/sq. cm
I	Hard brittle chalk with widely-spaced, tight joints	> 50 000	> 10	Negligible creep for pressures of at least 4 kg/sq. cm.

figure 2.3.4 (Dearman 1981) Correlation between engineering geological grade and mechanical properties of chalk at Mundford, England.

3 Sampling of Limestone

In order to be able to examine a wide range in mechanical and other properties it seemed most efficient to sample limestones from the various geological formations present in Europe. At the same time the relation between geological age and strength promised to be interesting. Accordingly samples were taken ranging from the limestone beds of the Silurian upto the Cretaceous calcarenites.

3.1 Sample Selection

During April 1987 twelve types of limestones were sampled from quarries in the U.K. ,Belgium and The Netherlands. Because for testing purposes solid fresh and homogeneous material was needed, large blocks of each type of limestone were selected from the highest quality zones in the quarries. Directly after this the blocks were wrapped in polyethylene bags in order to prevent the samples from drying out. In the case of the 'red marble' from Belgium it was not possible to acquire fresh material. Later in this report the test results indicate that the rock was moderately weathered.

Nearly all of the quarries in the U.K. used explosives to fragment the rockmass, while in Belgium and in Southern Limburg the limestone was cut with diamond wire or saws. Therefore it is possible that the blasting has induced fine cracks or extended preexisting cracks, which could influence the test results.

3.2 Sample Description

The description of the limestones was done following the IAEG report (1981) for engineering geological mapping (Appendix A.2). Furthermore the limestones are identified with local name, origin and geological age. The twelve samples were :

- Sample 01 The Magnesian Limestone from Yorkshire is of Permian age. It is described as a yellowish white medium grained fresh weak calcisiltite.
- Sample 02 The Chalk from near Hull is of Upper Cretaceous age, it is described as a pale white fine grained thinly laminated fresh weak calcisiltite.
- Sample 03 The Lower Oolite Limestone from Lincolnshire is of Jurassic age. It is described as a dark grey medium grained slightly weathered strong

limestone.

- Sample 04 The Upper Oolitic Limestone, also known as Corallian Limestone, from Yorkshire is of Jurassic age. It is described as a light grey medium grained fresh strong limestone.
- Sample 05 The Wenlock Limestone from Shropshire is of Silurian age. It is described as a light pinkish white coarse crystalline fossiliferous fresh strong limestone.
- Sample 06 Wooldale Limestone from Derbyshire is of Carboniferous age (Visean). It is described as a dark grayish black medium grained calcite veined fresh very strong limestone.
- Sample 07 Vinalmont Limestone, also known as Maassteen, from Belgium is of Carboniferous age (Visean). It is described as a light grey medium grained fresh very strong limestone.
- Sample 08 The Muschelkalk also known as Wellenkalk or wavy limestone from Gelderland is of Triassic age. It is described as a light whitish green fine grained thinly bedded fresh moderately strong micritic calcilutite.
- Sample 09 Belgian Fossil Limestone also known as Petit Granit, from the Dinant area in Belgium is of Carboniferous age (Tournaisian). It is described as a light grey medium grained fresh very strong limestone.
- Sample 10 Red Reef Marble from South Belgium is of Upper Devonian age (Frasnian). It is described as a whitish red fine crystalline fossiliferous moderately weathered strong limestone.
- Sample 11 Grey Marble from East Belgium is of Middle Devonian age (Givetian). It is described as a dark blackish grey medium size crystalline fossiliferous slightly weathered very strong limestone.
- Sample 12 Marl from South Limburg is of Upper Cretaceous age (Maastrichtian). It is described as a light yellowish beige medium grained fresh weak calcarenite.

An extensive engineering geological description of the rock

ENGINEERING GEOLOGICAL DESCRIPTION OF LIMESTONE
(According to I.A.E.G. Report for Description of Rocks)

CODES	AGE	ORIGIN	LOCATION	NAME	COORDIN. LONG./LAT.	GRAIN SIZE
SL01	PERMIAN	NEWTORPE YORKSHIRE	NEWTORPE QUARRY	MAGNESIAN LIMESTONE	W 1.17.. N 53.42..	MEDIUM
SL02	CRETACEOUS	ELSHAM YORKSHIRE	SINGLETON BIRCH QUARRY	UPPER CHALK	W 0.17.. N 53.32..	MEDIUM
SL03	JURASSIC	BRIGG LINCOLNSHIRE	LINDSEY QUARRY	LOWER OOLITE LST.	W 0.32.. N 53.24..	FINE
SL04	JURASSIC	PICKERING YORKSHIRE	HARGREAVES QUARRY	UPPER OOLITE LST.	W 0.47.. N 54.12..	COARSE
SL05	SILURIAN	MUCH WENLOCK WALES	SHADWELL QUARRY	WENLOCK LST.	W 2.34.. N 53.09..	MEDIUM
SL06	CARBONIFEROUS	BUXTON	TARMAC QUARRY	WOOLDALE LST.	W 1.53.. N 53.09..	MEDIUM
SL07	CARBONIFEROUS	VINALMONT	HAINAULT CARRIERE	VINALMONT LST.	O 4.32.. N 50.07..	MEDIUM
SL08	TRIASSIC	WINTERSWIJK GELDERLAND	ANKERSMIT GROEVE	MUSCHELKALK	O 5.55.. N 51.08..	FINE
SL09	CARBONIFEROUS	DENEE	MGP	TOURNASIAN LST.C3	O 4.45.. N 50.19..	FINE
SL10	DEVONIAN	NEUVILLE	NEUVILLE CARRIERE	MARBRE ROUGE	O 4.32.. N 50.07..	FINE
SL11	DEVONIAN	AYWAILLE	V/D WILDENBERG	MARBRE NOIR	O 5.41.. N 50.27..	FINE
SL12	CRETACEOUS	SIBBE LIMBURG	MERGELBOUW V/D KLEIN	MERGEL	O 5.49.. N 50.49..	MEDIUM

CODES	STRENGTH	JOINT SPACING	JOINT APERTURE	BEDDING	CEMENTATION	COLOUR
SL01	WEAK	MOD. WIDE	NARROW	VERY THICK	POORLY	LYW
SL02	MOD. WEAK	VERY WIDE	MEDIUM	VERY THICK	MODERATELY	LPW
SL03	STRONG	VERY WIDE	MEDIUM	VERY THICK	WELL	DBG
SL04	STRONG	VERY WIDE	OPEN	MEDIUM THICK	WELL	LYG
SL05	MOD. STRONG	MOD. WIDE	MEDIUM	MEDIUM THICK	WELL	LGW
SL06	STRONG	MOD. WIDE	OPEN	VERY THICK	MODERATELY	DGB
SL07	VERY STRONG	VERY WIDE	NARROW	VERY THICK	WELL	DBIG
SL08	MOD. WEAK	MOD. WIDE	MEDIUM	MEDIUM THICK	WELL	LBGr
SL09	VERY STRONG	VERY WIDE	NARROW	VERY THICK	WELL	LWG
SL10	STRONG	VERY WIDE	NARROW	VERY THICK	WELL	MWR
SL11	VERY STRONG	VERY WIDE	NARROW	VERY THICK	WELL	DGB
SL12	WEAK	VERY WIDE	NARROW	MEDIUM THICK	FRIABLE	LYB

CODES	WEATHERING	BEDDING TYPE	TEXTURE	ORIGINAL COMPONENTS	OTHER COMPONENTS
SL01	F	PARALLEL	GRANULAR	OOLITES	DOLOMITE
SL02	F	UNDULATING	GRANULAR	FORAMINIFERAE	FLINTS
SL03	S-M	PARALLEL	GRANULAR	PELLETS & MOLLUSCS	
SL04	S	PARALLEL	GRANULAR	OOLITES	SHELLS
SL05	S-M	UNDULATING	CRYSTALLINE	CORAL ORGANISMS	CALCITE CRYSTALS
SL06	S-M	PARALLEL	GRANULAR	CRINOIDS	CALCITE VEINS
SL07	F	PARALLEL	GRANULAR	OOLITES	MOLLUSCS
SL08	S	PARALLEL	GRANULAR	MUDS	PYRITE
SL09	F	PARALLEL	GRANULAR	CRINOIDS	CALCITE VEINS
SL10	M	UNDULATING	CRYSTALLINE	CORAL ORGANISMS	BRACHIOPODS
SL11	S	UNDULATING	CRYSTALLINE	CORAL ORGANISMS	BRACHIOPODS
SL12	F	PARALLEL	GRANULAR	PELLETS & SHELLS	FLINTS

FIGURE 3.2.1

masses from which the samples were taken is listed in fig. 3.2.1

3.3 Preparation of Specimens

A total of 300 cylindrical specimens were drilled for testing using core drilling bits. The number of test specimens of each limestone, subjected to one test, was six at maximum, provided the testing procedures required this. All samples were cored normal to the bedding and cored from a single block except for the Lower Oolite (SL03), the Wooldale Limestone (SL06) and the Muschelkalk (SL08). Diameters were 50, 40 and 30 mm with an maximum deviation of 0.1 % departure from the length axis.

With regard to the strength tests, 50 mm cores were taken from the weak limestones ; Magnesian Limestone (SL01), Upper Chalk (SL02), Muschelkalk (SL08) and Limburgian Marl (SL12). From the stronger limestones 40 mm specimens were cored, in order to ensure that the failure load did not exceed the maximum permissive load of the load frame. After coring surface grinding was applied to the 40 mm specimens for parallel planes. The 50 mm specimens could not endure such treatment without serious damage, due to their weakness. Finally the specimens were protected from deterioration by placing them in small airtight polyethylene bags.

Specimen failure during preparation sometimes occurred along cracks and calcite veins, as not uncommonly happens. This was especially the case with the calcite veined specimen of the Carboniferous Limestone SL06. A very sensitive sample to pre-testing failure was the Muschelkalk (SL08), due to its relatively high clay content and its thinly laminated bedding.

Tolerances on the dimensions of the cylindrical specimens were generally within the conditions defined in the procedures of the ASTM (American Society of Testing Materials) or the ISRM (International Society of Rock Mechanics).

Specimen No.	Material	Yield Point (ksi)	Tensile Strength (ksi)	Elongation (%)	Reduction of Area (%)	Impact (ft-lb)	Notes
101	Steel	45	70	25	60	10	
102	Steel	45	70	25	60	10	
103	Steel	45	70	25	60	10	
104	Steel	45	70	25	60	10	
105	Steel	45	70	25	60	10	
106	Steel	45	70	25	60	10	
107	Steel	45	70	25	60	10	
108	Steel	45	70	25	60	10	
109	Steel	45	70	25	60	10	
110	Steel	45	70	25	60	10	
111	Steel	45	70	25	60	10	
112	Steel	45	70	25	60	10	
113	Steel	45	70	25	60	10	
114	Steel	45	70	25	60	10	
115	Steel	45	70	25	60	10	
116	Steel	45	70	25	60	10	
117	Steel	45	70	25	60	10	
118	Steel	45	70	25	60	10	
119	Steel	45	70	25	60	10	
120	Steel	45	70	25	60	10	
121	Steel	45	70	25	60	10	
122	Steel	45	70	25	60	10	
123	Steel	45	70	25	60	10	
124	Steel	45	70	25	60	10	
125	Steel	45	70	25	60	10	
126	Steel	45	70	25	60	10	
127	Steel	45	70	25	60	10	
128	Steel	45	70	25	60	10	
129	Steel	45	70	25	60	10	
130	Steel	45	70	25	60	10	
131	Steel	45	70	25	60	10	
132	Steel	45	70	25	60	10	
133	Steel	45	70	25	60	10	
134	Steel	45	70	25	60	10	
135	Steel	45	70	25	60	10	
136	Steel	45	70	25	60	10	
137	Steel	45	70	25	60	10	
138	Steel	45	70	25	60	10	
139	Steel	45	70	25	60	10	
140	Steel	45	70	25	60	10	
141	Steel	45	70	25	60	10	
142	Steel	45	70	25	60	10	
143	Steel	45	70	25	60	10	
144	Steel	45	70	25	60	10	
145	Steel	45	70	25	60	10	
146	Steel	45	70	25	60	10	
147	Steel	45	70	25	60	10	
148	Steel	45	70	25	60	10	
149	Steel	45	70	25	60	10	
150	Steel	45	70	25	60	10	
151	Steel	45	70	25	60	10	
152	Steel	45	70	25	60	10	
153	Steel	45	70	25	60	10	
154	Steel	45	70	25	60	10	
155	Steel	45	70	25	60	10	
156	Steel	45	70	25	60	10	
157	Steel	45	70	25	60	10	
158	Steel	45	70	25	60	10	
159	Steel	45	70	25	60	10	
160	Steel	45	70	25	60	10	
161	Steel	45	70	25	60	10	
162	Steel	45	70	25	60	10	
163	Steel	45	70	25	60	10	
164	Steel	45	70	25	60	10	
165	Steel	45	70	25	60	10	
166	Steel	45	70	25	60	10	
167	Steel	45	70	25	60	10	
168	Steel	45	70	25	60	10	
169	Steel	45	70	25	60	10	
170	Steel	45	70	25	60	10	
171	Steel	45	70	25	60	10	
172	Steel	45	70	25	60	10	
173	Steel	45	70	25	60	10	
174	Steel	45	70	25	60	10	
175	Steel	45	70	25	60	10	
176	Steel	45	70	25	60	10	
177	Steel	45	70	25	60	10	
178	Steel	45	70	25	60	10	
179	Steel	45	70	25	60	10	
180	Steel	45	70	25	60	10	
181	Steel	45	70	25	60	10	
182	Steel	45	70	25	60	10	
183	Steel	45	70	25	60	10	
184	Steel	45	70	25	60	10	
185	Steel	45	70	25	60	10	
186	Steel	45	70	25	60	10	
187	Steel	45	70	25	60	10	
188	Steel	45	70	25	60	10	
189	Steel	45	70	25	60	10	
190	Steel	45	70	25	60	10	
191	Steel	45	70	25	60	10	
192	Steel	45	70	25	60	10	
193	Steel	45	70	25	60	10	
194	Steel	45	70	25	60	10	
195	Steel	45	70	25	60	10	
196	Steel	45	70	25	60	10	
197	Steel	45	70	25	60	10	
198	Steel	45	70	25	60	10	
199	Steel	45	70	25	60	10	
200	Steel	45	70	25	60	10	

4 Testing Techniques

Most tests were undertaken following ASTM or ISRM testing techniques. The sections following describe the most important features of the test methods and testing apparatus. The mechanical tests were performed on specimens at a moisture content measured after preparation of these test specimens. This is the moisture content given in table 4.11.4. Mechanical tests were performed normal to the bedding except in the case of the tensile strength. The tensile strength had to be tested parallel to the bedding because of the testing method. Hence a bedding description is necessary.

4.1 Water Content and Density ISRM suggested Method No. 2

The test was performed on three specimens with a diameter of 30 mm and a length/diameter ratio of 1:1. The tests involve determination of mass under field conditions as well as after drying in an oven at 105 degrees centigrade. The average mass of the cores was 50 grams. The accuracy to which the mass was recorded is + 0.005 grams and the dimensions of the cores were measured to the nearest 0.05 mm.

4.2 Porosity Method of Kobe

The test was undertaken following the method of Kobe with a "Ruska" universal porositymeter (appendix B.1). Oven dried cores of 30 mm diameter length/diameter ratio 1:1 were submerged in the pycnometer and their volume was measured without and with an internal pressure of 30 Psi (=206.8 KPa). The dimensions of the cores were measured to the nearest 0.05 mm and the accuracy of the volume measurements was + 0.005 ml.

4.3 Permeability

The permeability measurements were performed with a liquid-permeameter for the Limburgian Marl (SL02) cores and with a "Ruska" gaspermeameter for the other cores (appendix B.2). The liquid-permeameter uses water to determine the flowrate and is suitable for rather permeable media, such as this Limburgian Marl. The permeability using this apparatus is calculated following the formula:

4. Testing Techniques

Most tests were undertaken following ASTM or ISRM testing techniques. The sections following describe the most important features of the test methods and testing apparatus. The mechanical tests were performed on specimens of a material content measured after preparation of these test specimens. This is the material content given in Table 4.1.4. Mechanical tests were performed normal to the bedding except in the case of the tensile strength. The tensile strength had to be tested parallel to the bedding because of the testing method. Hence a bedding description is necessary.

4.1 Water Content and Density ISRM suggested Method No. 2

The test was performed on three specimens with a diameter of 30 mm and a length/diameter ratio of 1:1. The tests involve determination of mass under field conditions as well as after drying in an oven at 105 degrees centigrade. The average mass of the cores was 80 grams. The accuracy to which the mass was recorded is ± 0.005 grams and the dimensions of the cores were measured to the nearest 0.05 mm.

4.2 Porosity Method of Kober

The test was undertaken following the method of Kober with a "Rushka" universal porosimeter (Appendix B.1). Oven dried cores of 30 mm diameter length/diameter ratio 1:1 were submerged in the porosimeter and their volume was measured without and with an internal pressure of 30 Psi (206.8 KPa). The dimensions of the cores were measured to the nearest 0.05 mm and the accuracy of the volume measurements was ± 0.005 ml.

4.3 Permeability

The permeability measurements were performed with a liquid-permeameter for the "Rushka" (Appendix B.2) and with a "Rushka" gas-permeameter for the other cores (Appendix B.3). The liquid-permeameter uses water to determine the flow rate and is suitable for rather permeable media, such as this limestone. The permeability using this apparatus is calculated following the formula:

$$K = \frac{m L V}{A dP t}$$

where m = viscosity (Cp) ; $m_{\text{water}} = 1 \text{ Cp}$ (Cp = centi poise)
 L = length of the sample (mm)
 V = volume passed (cc)
 A = area normal to the flow (mm)
 dP = difference in pressure (KPa)
 t = time interval (sec)

The gaspermeameter allows for natural gas under a certain pressure to flow through the specimen in small quantities, hence the principle is the same as for the liquidpermeameter. This apparatus measures in the range of 10^{-3} Darcy.

The specimen of 30 mm diameter and length/diameter ratio of 1:1 were oven dried and their permeability was measured with a deviation of approximately 20 % due to turbulence and gas solution in the water.

4.4 Tensile strength ASTM standard D 3967-81

The tensile strength was determined with an indirect method known as the Brazilian Tensile Strength test (BTS test). In the procedure of this test discs of with a 50 mm diameter and 25 mm thickness are split by vertical loading through a diameter.

In the Wykeham-Farrance 100 kN strain guided loading frame platens were connected to LVDT (Linear Variable Differential Transducers) displacement transducers. The LVDT's used are electromechanical transformers with a range of + 1 mm. The load was transmitted by a 50 kN load transducer. Through an 12 bits A/D converter displacements and load were digitized and recorded for every 0.001 mm vertical deformation. The displacement rate of the loading frame was 0.1 mm/min.

4.5 Unconfined Double Shear Test

This test determines the direct shear strength of intact rock as suggested by T.R. Stacey (1980). The test frame was developed by the section of Engineering Geology of the Technical University of Delft. The method involves punching a steel block trough a rock disc of 50 mm diameter and a Thickness of 10 mm .Two predetermined shear planes show when the failure occurs. Loading should be applied at a rate of 15 kN/min.

The shear test apparatus was placed in the ELE point load testing frame. The Unconfined Shear Strength was calculated with

$$k = \frac{V \cdot L}{A \cdot \Delta P \cdot t}$$

where k = viscosity (cP); V = volume passed (cc); L = length of the sample (mm); A = area normal to the flow (mm²); ΔP = difference in pressure (KPa); t = time interval (sec).

The viscometer allows for natural gas under a certain pressure to flow through the specimen in small quantities, hence the principle is the same as for the liquid viscometer. This apparatus measures in the range of 10⁻³ cP.

The specimen of 30 mm diameter and length/diameter ratio of 1:1 were oven dried and their permeability was measured with a deviation of approximately 20% due to turbulence and gas solution in the water.

4.4 Tensile Strength ASTM standard D 3987-81

The tensile strength was determined with an indirect method known as the Brazilian Tensile Strength Test (ASTM Test). In the procedure of this test discs of 50 mm diameter and 25 mm thickness are split by vertical loading through a diameter.

In the Wykeham-Farrance 100 kN strain guided loading frame platens were connected to LVDT (Linear Variable Differential Transducer) displacement transducers. The LVDT's used are electro-mechanical transducers with a range of +1 mm. The load was transmitted by a 50 kN load transducer. Through an 18 bit A/D converter displacements and load were digitized and recorded for every 0.001 mm vertical deformation. The displacement rate of the loading frame was 0.1 mm/min.

4.5 Unconfined Simple Shear Test

This test determines the direct shear strength of intact rock as suggested by F.H. Kreyer (1986). The test frame was developed by the section of Engineering Geology of the Technical University of Delft. The method involves punching a steel block through a rock disc of 50 mm diameter and a thickness of 10 mm. Two predetermined shear planes where the failure occurs. Loading should be applied at a rate of 10 kN/min.

The shear test apparatus was placed in the ELS point load testing frame. The Unconfined Shear Strength was calculated with

the formula :

$$USS = \frac{P}{T * (L_1 + L_2)}$$

USS = Unconfined Shear Strength (MPa)

P = failure load (kN)

L₁&L₂ = length's of the shear planes (mm)

T = disc thickness (mm)

Because no clear planes developed the measurement of the shear planes in some cases had to be estimated. The accuracy of the length measurement was + 0.05 mm, while the load gauges deviated approximately 1 kN .

4.6 Unconfined Compressive Strength, Elastic-modulus and Poisson's ratio ASTM D 2938-79, ASTM 3148-80

The Unconfined Compressive Strength (UCS) was determined using a 700 kN closed loop servo controlled load frame (see appendix B.3) for the 40 mm length cores and the WF 100 kN load frame for the 50 mm length cores. The length diameter ratio of the test specimens was 2:1 . The rate of loading was adapted so that the time in which the sample failed was 5 to 10 minutes. The exact loading rate range was 5.6 - 11.9 MPa/min.

Using LVDT transducers the vertical displacements were twice measured in opposite directions. The lateral displacements were four times measured at angles of 90 degrees. The measurement interval was fixed at 0.001 mm vertical deformation. This implied that for every 0.001 mm vertical deformation data were stored in the computer. Together with the signal from the load transducers they were digitized and converted to mean values for the proper stress-strain relations which were recorded on floppy disk.

Then from the deformation curves the elastic moduli were calculated with a tangent method at 50 % of the failure stress or, if the slope at 50 % of the failure stress was not representative, the average was taken at the point of inflexion. At the same point the Poisson's ratio's were calculated, dividing the lateral strain by the vertical strain.

the formula:

$$\sigma_{ss} = \frac{F}{T \cdot (L_1 + L_2)}$$

σ_{ss} = Unconfined Shear Stress (MPa)
 F = Failure Load (kN)
 L_1 = length of the shear planes (mm)
 T = disc thickness (mm)

Because no clear planes developed the measurement of the shear planes in some cases had to be estimated. The accuracy of the length measurement was ± 0.05 mm, while the load gauges deviated approximately 1 kN.

4.3 Unconfined Compressive Strength, Elastic Modulus and Poisson's ratio
 ASTM D 2938-78, ASTM 318-80

The Unconfined Compressive Strength (UCS) was determined using a 700 kN closed loop servo controlled load frame (see appendix B.3). For the 40 mm length cores and the 50 mm length cores the length diameter ratio of the test specimens was 2:1. The rate of loading was adapted so that the time in which the sample failed was 5 to 10 minutes. The exact loading rate range was 0.5 - 11.5 MPa/min.

Using LVDT transducers the vertical displacements were measured in opposite directions. The lateral displacements were four times measured at angles of 90 degrees. The measurement interval was fixed at 0.001 mm vertical deformation. This implied that for every 0.001 mm vertical deformation data were stored in the computer. Together with the signal from the load transducers they were digitized and converted to mean values for the proper stress-strain relations which were recorded on floppy disk.

Then from the deformation curves the elastic modulus was calculated with a tangent method at 50 % of the failure stress or at the slope of 50 % of the failure stress was not representative, the average was taken at the point of inflection. At the same point the Poisson's ratio's were calculated, dividing the lateral strain by the vertical strain.

4.7 Ultra Sonic Velocity

ISRM suggested method No. 4

This index test was undertaken on the specimens prepared for the UCS using a CCT 4 ultra sonic velocity concrete tester. Pulses are generated by a 10 MHz oscillator and passed through the sample. Travel times are measured continuously in microseconds. To exclude the signal from other media a contact fluid is applied to both ends of the core.

4.8 Slake Durability

ISRM suggested document 2 part 2

The slake durability test is a combination of abrasion and slaking performed on 10 lumps of the sample. The limestone lumps are agitated by revolving them in a cylindrical mesh drum in cycles of 10 minutes, immersed in water. Following this the retained material in the drum is oven dried at 105 degrees centigrade and weighed. The Slake Durability Index is then calculated from the formula given below:

$$Id_2 = \frac{W_3}{W_1} * 100 \%$$

were : Id_2 = slake durability index

W_3 = weight after the second cycle

W_1 = weight before first cycle

The accuracy of the balance measurements + 0.005 grams.

4.9 X-Ray Fluorescence

The percentages $CaCO_3$ and MgO were determined with the X-Ray fluorescence technique from the Rontgen Laboratory of the Department of Mining and Petroleum Engineering (appendix B.4).

Sample preparation was performed by crushing representative core specimen to maximum diameter of 0.02 mm using a disc-crusher. Next the crushed sample was mixed and divided into equal portions.

The analyses were conducted on glass pearls (0.5 g sample + 5.0 g $Li_2B_4O_7$). Thus the pearls were compared with standard samples of the same matrix. With the aid of a linear regression method the percentages were calculated. Thereupon corrections for inter-element effects and weighing inaccuracy were applied. The total error of the calculated values is no more than 0.5 % .

4.7 Ultrasonic Velocity ISRM suggested method No. 4

This index test was undertaken on the specimens prepared for the UCS using a GTS 4 ultrasonic velocity concrete tester. Pulses are generated by a 10 MHz oscillator and passed through the sample. Travel time are measured continuously in microseconds. To exclude the signal from other media a contact fluid is applied to both ends of the core.

4.8 Slake Durability ISRM suggested document 2 part 2

The slake durability test is a combination of abrasion and slaking performed on 10 jumps of the sample. The slake jumps are initiated by revolving them in a cylindrical wash drum in cycles of 10 minutes, immersed in water. Following this the retained material in the drum is oven dried at 105 degrees centigrade and weighed. The Slake Durability Index is then calculated from the formula given below:

$$Id_s = \frac{W_2}{W_1} \times 100 \%$$

where: Id_s = slake durability index
 W_2 = weight after the second cycle
 W_1 = weight before first cycle
The accuracy of the balance measurement is 0.005 grams.

4.9 X-Ray Fluorescence

The percentages CaO and MgO were determined with the X-Ray fluorescence technique from the Rockwell Laboratory of the Department of Mining and Petroleum Engineering (Appendix B.4).

Sample preparation was performed by crushing representative core specimens to maximum diameter of 0.85 mm using a disc crusher. Next the crushed sample was mixed and divided into equal portions.

The analyses were conducted on glass beads (0.5 g sample + 0.5 g LiBrO₂). Thus the beads were separated with standard samples of the same matrix. With the aid of a linear regression method the percentages were calculated. Thereupon corrections for inter-element effects and weighing inaccuracy were applied. The total error of the calculated values is no more than 0.5 %.

4.10 Carbonate Resistance Index (CRI)

This test was developed by the author with the purpose to give an indication of the relative solubility rate of the carbonate rocks. The test may be considered valid only for those limestones, consisting for the major part of calcium and/or magnesium carbonate and in which no other significant in acid soluble minerals occur.

The test involves the submerging of a dried and weighed test specimen (the one used was cylindrical) in a 1.0 N hydrochloric acid. The specimen remains in 150 ml of this solution for exact 5 minutes after the specimen is oven dried and weighed. The CRI is calculated with the following formula :

$$CRI = \frac{\text{Loss of weight}}{\text{Original weight}} * \frac{\text{Surface area of the specimen}}{\text{Surface area of a sphere with the same volume}} * (\text{CO}_3\text{-correction})$$

when applied to a cylindrical specimen the CR-Index is :

$$CRI = \frac{(W_1 - W_2)}{W_1} \cdot \frac{(2\pi \cdot R_d + 2\pi \cdot R_d \cdot D)}{(4\pi \cdot R_s^2)} \cdot \frac{10000}{\%CO_3}$$

were : W_1 = dry weight before testing
 W_2 = dry weight after testing
 R_d = diameter of the cylinder
 R_s = diameter of imaginary sphere of the same content
 D = thickness of the cylinder
 $\% CO_3$ = total amount of carbonate material

R_s is calculated from the formula : $R_s = \sqrt[3]{\{ 3/4 \cdot R_d \cdot D \}}$

is represented by the formula: $\alpha = \sqrt{\frac{2 \cdot H \cdot W}{\pi \cdot D}}$

where: α - angle of contact between
 D - diameter of the cylinder
 H - diameter of (horizontal) plate of the same material
 W - diameter of the cylinder
 M - all weight of the contact
 m - all weight of the contact

$$CHI = \frac{M}{(M - m)} \cdot \frac{(2 \cdot H \cdot W)}{(2 \cdot H \cdot W + 2 \cdot H \cdot W \cdot D)} \cdot \frac{2000}{10000}$$

where according to a simplified definition the CB-index is:

$$CHI = \frac{\text{original weight of the specimen}}{\text{weight of the specimen}} \cdot \frac{\text{surface area of the specimen}}{\text{surface area of the specimen}} \cdot (C00 - collection)$$

CHI is calculated with the following formula:
 where: α - angle of contact between
 D - diameter of the cylinder
 H - diameter of (horizontal) plate of the same material
 W - diameter of the cylinder
 M - all weight of the contact
 m - all weight of the contact

where: α - angle of contact between
 D - diameter of the cylinder
 H - diameter of (horizontal) plate of the same material
 W - diameter of the cylinder
 M - all weight of the contact
 m - all weight of the contact

where: α - angle of contact between (CHI)

The temperature at which the test was performed was 22 degrees centigrade. For each test a new volume of acid is required. The measurement accuracy is + 0.05 mm. for length's and + 0.005 gram for weight.

4.11. Presentation of the Test Results

The complete record of all calculated test results is given in tables 4.11.2 - 4.11.4 . In these tables the above mentioned parameters are indicated for the twelve limestone samples. To simplify the presentation codes for the samples and abbreviations for the parameters, which are explained at the top of the table, are used (table 4.11.1). The accuracies are conform the limitations of the performed tests.

4.12 Microscope Study

The main aim of the microscope study was the identification of features related to engineering properties. As mentioned in chapter 2 the important contaminant minerals, besides carbonates, can be quartz, both detrital and silica quartz, and clay minerals such as bentonite and illite. Small amounts of carbonaceous fines cause the grey colour in the limestones. From physical point of view the grain sizes and the hardnesses of the various minerals form a point of investigation. The cuttability and the abrasive capacity are engineering parameters directly related to these factors.

From the limestone samples 12 thin sections have been examined under a transmitted light polarizing microscope. With the use of density diagrams the percentages of non-carbonate minerals have been estimated. The determination of grain sizes was done following the linear intercept method ; here the average intercept length has been multiplied by a correction factor of 1.5 as suggested in DIN 22021 (1985).

The data are summarized in table 4.12.1 where also the origin of the components is given. If the rock consists of particles and matrix, the grain size of both is indicated.

The temperature at which the test was performed was 25 degrees centigrade. For each test a new volume of acid is required. The measurement accuracy is ± 0.05 mm. for length and ± 0.005 gram for weight.

4.11 Presentation of the Test Results

The complete record of all calculated test results is given in Tables 4.11.1 - 4.11.4. In these tables the above mentioned parameters are indicated for the twelve limestone samples. To simplify the presentation codes for the samples and abbreviations for the parameters, which are explained at the top of the tables, are used (Table 4.11.1). The notations are chosen to conform the limitations of the performed tests.

4.12 Microscope Study

The main aim of the microscope study was the identification of features related to engineering properties. As mentioned in Chapter 2 the important constituent minerals, besides carbonates, can be quartz, both detrital and siliceous quartz, and clay minerals such as bentonite and illite. Small amounts of carbonaceous fines cause the grey color in the limestones. From physical point of view the grain sizes and the hardnesses of the various minerals form a point of investigation. The cutability and the abrasive capacity are engineering parameters directly related to these factors.

From the limestone samples 12 thin sections have been examined under a transmitted light polarizing microscope. With the use of density diagrams the percentages of non-carbonate minerals have been estimated. The determination of grain sizes was done following the linear intercept method; here the average intercept length has been multiplied by a correction factor of 1.5 as suggested in DIN 52021 (1982).

The data are summarized in Table 4.12.1 where also the origin of the components is given. If the rock consists of particles and matrix, the grain size of both is indicated.

LEGEND for TEST RESULTS

TYPE OF TEST :

UCS = Unconfined Compressive Strength (MPa)
BTS = Brazilian Tensile Strength (MPa)
USS = Unconfined Shear Strength (MPa)
E-Modulus = Deformation Constant (GPa)
Poisson's Ratio = Lateral/Vertical Strain
Velocity = Ultra Sonic Velocity (m/s)
Porosity = Effective Porosity (%)
Permeab. = Permeability (mDarcy)
N.M.Cont = Natural Moisture Content (%)
Dry Density = Dry density (Tons/M³)
Nat. Density = Natural Density (Tons/M³)
CR-Index = Carbonate Resistance Index
Id2 =Slake Durability Index
MgCO₃ % = Magnesium Carbonate % (Dolomite)
CaCO₃ % = Calcium Carbonate %
XRF ANALYSIS = X-Ray Diffraction

STATISTIC CODES (see table 5.1.1)

Nx = Number of test results
[S.D.] = Standard Deviation
Avg. = Average Value
* for median average values of tested sets
+ for mean average values of tested sets

TYPE OF LIMESTONE :

SL01 = Magnesian Limestone
SL02 = Upper Chalk
SL03 = Lower Dolite
SL04 = Upper Dolite
SL05 = Wenlock Limestone
SL06 = Carboniferous Limestone
SL07 = Maassteen (Crinoid Lst.)
SL08 = Muschelkalk
SL09 = Belgian Fossil Limestone
SL10 = Red Marble
SL11 = Grey Marble
SL12 = Limburgian Marl

SAMPLE CODES (see tables 4.11.1-3)

For Example SL03.2.1 means :

SL = Sample
03 = Third sample type
.2 = Second test series
.1 = First specimen

SAMPLE CODES	U.C.S. MPa	E-MODULUS GPa	POISSON RATIO	VELOCITY M/S	SAMPLE CODES	BTS MPa	STRAIN 10 ⁻³ mm	SAMPLE CODES	USS MPa
SL01.1.1	11.69	2.88	.45	2421	SL01.2.1	1.38	175.42	SL01.3.1	7.92
SL01.1.2	10.18	4.22	.30	2509	SL01.2.2	2.41	223.02	SL01.3.2	7.76
SL01.1.3	13.23	5.35	.19	2612	SL01.2.3	2.15	163.08	SL01.3.3	4.47
SL01.1.4	13.73	4.71	.31	2604	SL01.2.4	1.03	161.50	SL01.3.4	5.21
SL01.1.5	15.29	5.27	.13	2603	SL01.2.5	2.22	3.91	SL01.3.5	4.45
SL01.1.6	12.68	4.24	.68	2750	SL01.2.6	2.18	197.87	SL01.3.6	7.02
SL02.1.1	24.89	10.90	.16	4297	SL02.2.1	3.23	158.44	SL02.3.1	13.78
SL02.1.2	13.48	8.91	.15	4197	SL02.2.2	2.38	345.21	SL02.3.2	11.27
SL02.1.3	26.73	10.99	.25	4233	SL02.2.3	3.37	186.65	SL02.3.3	16.57
SL02.1.4	19.98	9.33	.11	4111	SL02.2.4	1.53	151.61	SL02.3.4	14.17
SL02.1.5	24.63	13.39	.36	4416	SL02.2.5	3.43	167.96	SL02.3.5	9.80
SL02.1.6	25.27	13.39	.37	4057	SL02.2.6	3.74	157.10	SL02.3.6	13.34
SL03.1.1	96.07	40.35	.33	5570	SL03.2.1	6.87	154.54	SL03.3.1	22.04
SL03.1.2	86.75	39.89	.31	5594	SL03.2.2	5.64	127.32	SL03.3.2	20.91
SL03.1.3	115.79	38.49	.33	5575	SL03.2.3	6.72	96.56	SL03.3.3	25.14
SL03.1.4	92.67	40.20	.27	5525	SL03.2.4	6.05	95.46	SL03.3.4	22.65
SL03.1.5	116.47	41.19	.29	5564	SL03.2.5	6.21	103.89	SL03.3.5	14.45
SL03.1.6	95.00	42.47	.31	5534	SL03.2.6	5.93	82.03	SL03.3.6	15.26
SL04.1.1	138.52	37.37	.32	5319	SL04.2.1	5.62	135.25	SL04.3.1	21.61
SL04.1.2	117.25	30.96	.27	4990	SL04.2.2	6.08	186.09	SL04.3.2	20.26
SL04.1.3	125.80	31.16	.29	5103	SL04.2.3	6.88	135.86	SL04.3.3	16.31
SL04.1.4	80.72	29.86	.37	5025	SL04.2.4	6.81	110.10	SL04.3.4	19.80
SL04.1.5				5252	SL04.2.5	7.41	113.16	SL04.3.5	17.22
SL04.1.6	122.59	32.92	.31	5236	SL04.2.6	8.47	142.60	SL04.3.6	22.62
SL05.1.1	81.02	62.42	.33	5534	SL05.2.1	6.10	59.20	SL05.3.1	21.52
SL05.1.2	71.40	62.57	.33	5536	SL05.2.2	5.05	58.72	SL05.3.2	21.72
SL05.1.3	74.99	57.74	.39	5308	SL05.2.3	6.59	70.07	SL05.3.3	19.26
SL05.1.4	75.48	62.93	.36	5571	SL05.2.4	5.43	56.88	SL05.3.4	10.39
SL05.1.5	72.08	63.16	.36	5525	SL05.2.5	6.37	66.04	SL05.3.5	26.78
SL05.1.6	78.10	64.56	.39	5881	SL05.2.6	6.42	64.58	SL05.3.6	20.36
SL06.1.1				6277	SL06.2.1	9.31	90.33	SL06.3.1	22.33
SL06.1.2	107.83	64.92	.32	6455	SL06.2.2	7.51	72.63	SL06.3.2	19.88
SL06.1.3	56.83	62.38	.30	6372	SL06.2.3	9.53	84.96	SL06.3.3	24.40
SL06.1.4	86.07	62.62	.30	6404	SL06.2.4	7.59	65.06	SL06.3.4	22.64
SL06.1.5	170.87	64.94	.35	6285	SL06.2.5	8.01	80.08	SL06.3.5	30.26
SL06.1.6					SL06.2.6	4.53		SL06.3.6	24.70

SAMPLE CODES	U.C.S. MPa	E-MODULUS GPa	POISSON RATIO	VELOCITY M/S	SAMPLE CODES	BTS MPa	STRAIN 10 ⁻³ mm	SAMPLE CODES	USS MPa
SL07.1.1	195.06	70.09	.32	6569	SL07.2.1	6.57	61.17	SL07.3.1	32.70
SL07.1.2	185.83	69.94	.31	6526	SL07.2.2	7.90	83.84	SL07.3.2	24.36
SL07.1.3	190.78	70.18	.31	6562	SL07.2.3	8.14	75.81	SL07.3.3	24.57
SL07.1.4	191.85	70.83	.30	6569	SL07.2.4	7.70	76.66	SL07.3.4	25.41
SL07.1.5	194.67	70.37	.32	6559	SL07.2.5	6.91	73.12	SL07.3.5	22.79
SL07.1.6	184.37	70.61	.32	6536	SL07.2.6	6.84	74.71	SL07.3.6	-----
SL08.1.1	83.81	17.16	.37	4136	SL08.2.1	5.22	119.02	SL08.3.1	21.60
SL08.1.2	51.10	12.20	.32	4141	SL08.2.2	7.39	145.26	SL08.3.2	19.85
SL08.1.3	37.61	11.81	.70	4103	SL08.2.3	6.49	134.40	SL08.3.3	19.46
SL08.1.4	66.58	15.81	.44	4132	SL08.2.4	4.20	226.07	SL08.3.4	20.44
SL08.1.5	39.29	13.17	.19	4212	SL08.2.5	5.84	151.00	SL08.3.5	13.97
SL08.1.6	66.21	16.11	.27	4128	SL08.2.6	5.59	136.35	SL08.3.6	18.76
SL09.1.1	148.82	69.18	.29	6439	SL09.2.1	9.63	96.44	SL09.3.1	20.64
SL09.1.2	159.51	70.80	.34	6445	SL09.2.2	6.34	86.55	SL09.3.2	19.88
SL09.1.3	153.00	69.29	.32	6445	SL09.2.3	9.71	92.16	SL09.3.3	21.65
SL09.1.4	159.99	69.83	.34	6468	SL09.2.4	8.75	89.48	SL09.3.4	22.13
SL09.1.5	158.63	69.72	.35	6432	SL09.2.5	6.20	93.63	SL09.3.5	22.79
SL09.1.6	162.52	70.72	.35	6413	SL09.2.6	9.50	105.35	SL09.3.6	22.69
SL10.1.1	96.07	51.40	.22	6006	SL10.2.1	6.59	177.60	SL10.3.1	28.81
SL10.1.2	80.53	52.55	.34	6104	SL10.2.2	7.05	96.56	SL10.3.2	21.52
SL10.1.3	86.55	57.66	.27	6166	SL10.2.3	5.65	104.25	SL10.3.3	34.66
SL10.1.4	107.15	54.22	.28	6153	SL10.2.4	6.46	152.59	SL10.3.4	26.99
SL10.1.5	61.10	55.46	.37	6196	SL10.2.5	7.11	81.79	SL10.3.5	29.67
SL10.1.6	105.20	59.42	.32	6210	SL10.2.6	5.01	75.93	SL10.3.6	29.55
SL11.1.1	141.34	68.46	.32	6155	SL11.2.1	9.42	95.33	SL11.3.1	20.77
SL11.1.2	160.96	67.52	.29	6226	SL11.2.2	8.65	86.91	SL11.3.2	24.44
SL11.1.3	156.01	62.10	.30	6206	SL11.2.3	8.39	78.98	SL11.3.3	20.93
SL11.1.4	156.01	62.36	.30	6097	SL11.2.4	7.83	89.36	SL11.3.4	20.49
SL11.1.5	178.45	66.77	.34	6212	SL11.2.5	9.75	94.36	SL11.3.5	18.15
SL11.1.6	152.12	76.58	.34	6200	SL11.2.6	10.47	101.20	SL11.3.6	21.85
SL12.1.1	3.54	1.24	.22	1755	SL12.2.1	.41	430.78	SL12.3.1	1.07
SL12.1.2	3.72	1.29	.25	1746	SL12.2.2	.41	467.65	SL12.3.2	.68
SL12.1.3	4.11	1.17	.29	1741	SL12.2.3	.40	425.70	SL12.3.3	.59
SL12.1.4	3.46	1.25	.25	1792	SL12.2.4	.38	505.16	SL12.3.4	.94
SL12.1.5	2.86	1.01	.19	1772	SL12.2.5	.33	384.03	SL12.3.5	.69
SL12.1.6	3.41	1.28	.30	1790	SL12.2.6	.37	363.16	SL12.3.6	1.01

table 4.11.3

CODE SAMPLE	POROSITY (%)	PERMEABILITY m Darcy	DRY DENSITY (TON/M3)	NAT. DENSITY (TON/M3)	M.N.CONT. (%)	CR-INDEX	SLAKE DURA- BILITY Id2	XRF-ANALYSIS
SL01.4.1	38.16	2.46	1.75	2.11	17.38	154.3	80.91	CaCO3%-62.18
SL01.4.2	37.93	2.60	1.76	2.11	16.78	163.1		MgCO3%-37.92
SL01.4.3	37.81	3.57	1.77	2.13	16.95			
SL02.4.1	14.97	0	2.30	2.41	4.75	70.6	98.03	CaCO3%-97.60
SL02.4.2	16.59	0	2.25	2.38	5.26	75.0		MgCO3%- 0.90
SL02.4.3	15.53	0	2.28	2.40	5.23			
SL03.4.1	5.41	0	2.57	2.61	1.60	86.1	98.78	CaCO3%-92.66
SL03.4.2	2.71	0	2.59	2.63	1.36	85.5		MgCO3%- 1.19
SL03.4.3	3.27	0	2.58	2.62	1.50			
SL04.4.1	2.12	0	2.53	2.58	2.07	75.7	99.65	CaCO3%-83.64
SL04.4.2	2.09	0	2.55	2.59	1.65	80.6		MgCO3%- 0.99
SL04.4.3	1.79	0	2.51	2.57	2.11			
SL05.4.1	1.11	0	2.68	2.68	.02	79.3	98.86	CaCO3%-97.41
SL05.4.2	1.14	0	2.68	2.68	.04	82.1		MgCO3%- 1.10
SL05.4.3	.98	0	2.68	2.68	.02			
SL06.4.1	.60	0	2.67	2.67	.19	75.0	98.11	CaCO3%-98.40
SL06.4.2	.49	0	2.68	2.68	.13	78.7		MgCO3%- 1.13
SL06.4.3	.61	0	2.67	2.68	.21			
SL07.4.1	.37	0	2.69	2.69	.05	88.8	99.32	CaCO3%-99.37
SL07.4.2	.70	0	2.68	2.68	.09	91.1		MgCO3%- 0.74
SL07.4.3	.66	0	2.68	2.69	.09			
SL08.4.1	6.13	0	2.56	2.58	.69	108.6	96.31	CaCO3%-82.01
SL08.4.2	4.02	0	2.55	2.57	.64	113.6		MgCO3%- 4.29
SL08.4.3	6.10	0	2.54	2.56	.79			
SL09.4.1	.88	0	2.67	2.67	.02	78.2	99.38	CaCO3%-97.13
SL09.4.2	1.04	0	2.66	2.66	.03	78.8		MgCO3%- 1.51
SL09.4.3	.99	0	2.67	2.67	.05			
SL10.4.1	.64	0	2.69	2.69	.07	75.9	98.99	CaCO3%-91.00
SL10.4.2	.63	0	2.69	2.69	.13	78.4		MgCO3%- 2.00
SL10.4.3	.49	0	2.69	2.70	.05			
SL11.4.1	.84	0	2.68	2.69	.19	75.5	99.48	CaCO3%-96.71
SL11.4.2	.76	0	2.68	2.69	.15	77.2		MgCO3%- 1.99
SL11.4.3	.78	0	2.68	2.68	.16			
SL12.4.1	50.15	4600	1.34	1.72	22.20	258.0	71.41	CaCO3%-96.64
SL12.4.2	50.02	5500	1.34	1.71	21.98	266.6		MgCO3%- 1.70
SL12.4.3	49.97	5400	1.33	1.71	21.97			

table 4.11.4

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MICROSCOPE STUDY OF THIN SECTIONS

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SAMPLE CODES	GRAINSIZE mm		QUARTZ (%)	FELDSPAR (%)	**ARG.FINES (%)	CARBON	SHAPE OF PART./CRYST.	TYPE OF COMPONENT	ABRASIVE CAPACITY
SL01	.13	Particle					Rounded	Oolitic	0
	.001	Matrix							
SL02	.07	Particle	+		+		Rounded	Skeletal	1
	.002	Matrix							
SL03	.80	Particle	++	+		+	Sub Rounded	Pellets	2
	.07	Matrix							
SL04	.09	Particle			00	+	Sub Rounded	Sketetal	1
	.002	Matrix						+ Muds	
SL05	4.80	Crystals		+			Irregular	Skeletal (Reefoidal)	1
SL06	.09	Particle				+	Rounded	Skeletal	1
		*Crys.cem.							
SL07	.48	Particle				+	Rounded	Skeletal	2
		*Crys.cem.						(Crinoidal)	
SL08	.01	Particle			00		Sub Angular	Muds	1
		*Crys.cem.							
SL09	.60	Particle	+			+	Sub Angular	Oolitic	2
		*Crys.cem.						+ Fossils	
SL10	.01	Crystals	+	+			Angular	Skeletal	2
								(Reefoidal)	
SL11	.08	Crystals		+		+	Angular	Skeletal	2
								(Reefoidal)	
SL12	.12	Particle					Rounded	Pellets	0
	.01	Matrix							

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LEGEND OF MICROSCOPE DATA

MODAL VOLUMES OF MINERALS :

00 = 6-10 %
 0 = 3-6 %
 ++ = 1-3 %
 + = < 1 %

ABRASIVE CAPACITY :

F = Quartz % * Grainsize *
 BTS * 10

CUTTABILITY :

Ratio (UCS/BTS)

TYPE OF LIMESTONE :

SL01 = Magnesian Limestone
 SL02 = Upper Chalk
 SL03 = Lower Oolite
 SL04 = Upper Oolite
 SL05 = Wenlock Limestone
 SL06 = Carboniferous Limestone
 SL07 = Maassteen (Crinoid Lst.)
 SL08 = Muschelkalk
 SL09 = Belgian Fossil Limestone
 SL10 = Red Marble
 SL11 = Grey Marble
 SL12 = Limburgian Marl

* = Crystalline Cement
 *** = Argillaceous Fines

5 Evaluation of test results

In order to be able to rely upon the information about the test results, the strength's and weaknesses of the various data must be assessed. This can be done with the help of statistics and a knowledge of the limestones in relation to their properties. When the test parameters are also compared it is possible say something about the degree of correlation.

5.1 Statistical Analysis

Distributions of the performed tests have certain characteristics such as their midpoint; measures indicating their spread; and measures of symmetry of the distribution. These characteristics are known as parameters if they describe the population of a limestone, and statistics if they refer to sets of limestone test data. In the following sections the statistics are used to estimate the parameters of the parent population, which is represented by the single block, and to test hypotheses about these populations.

The most obvious of a data set is some type of average value. Two of the important ones are the mean and the median; the mean representing the sum of the data divided by the number of data; and the median being the value, which at ranking in height takes the most middlemost value. To decide which average renders the most realistic value for the limestone properties concerned, the character of the frequency distribution is of importance. In a normal distribution, which is symmetric, the highest frequency coincides with the mean and the median. Since in this case the mean uses the values of all the data, and is demonstrated to be closer to the population mean than any other average, it is the most useful estimator. If however the distribution is strongly asymmetric the median may render more information about the population of limestone properties, because it is not so sensitive to single values with a large deviation. In strength properties the median value tends to be higher than the mean value. This is caused due to the fact that local weak planes in the block of limestone result in low test values, which are by no means representative for the rock material.

A set of data consists of all the values related to one test parameter and to one type of limestone (table 4.11.1). To test the hypothesis whether a data set is normal or not, and whether the mean value should be used or not, only the sets of 6 data can be subjected to such a normality test. Concerning the sets with less data, their number is too small to predict their distribution. Hence only a mean value can be calculated.

i	x(i)	(x(i) - \bar{x})/s	$t_i = \hat{F}(x(i))$	$(t_i - \frac{2i-1}{2n})^2$
1	1,34	-1,581	0,0569	0,00005
2	1,38	-1,186	0,1178	0,00104
3	1,41	-0,889	0,1870	0,00397
4	1,47	-0,296	0,3836	0,00113
5	1,49	-0,099	0,4606	0,00011
6	1,53	0,296	0,6164	0,00441
7	1,56	0,593	0,7234	0,00539
8	1,57	0,692	0,7555	0,00003
9	1,59	0,889	0,8130	0,00137
10	1,66	1,581	0,9431	0,00005
$W^2 = 0,01755 + \frac{1}{120} = 0,02588$				0,01755

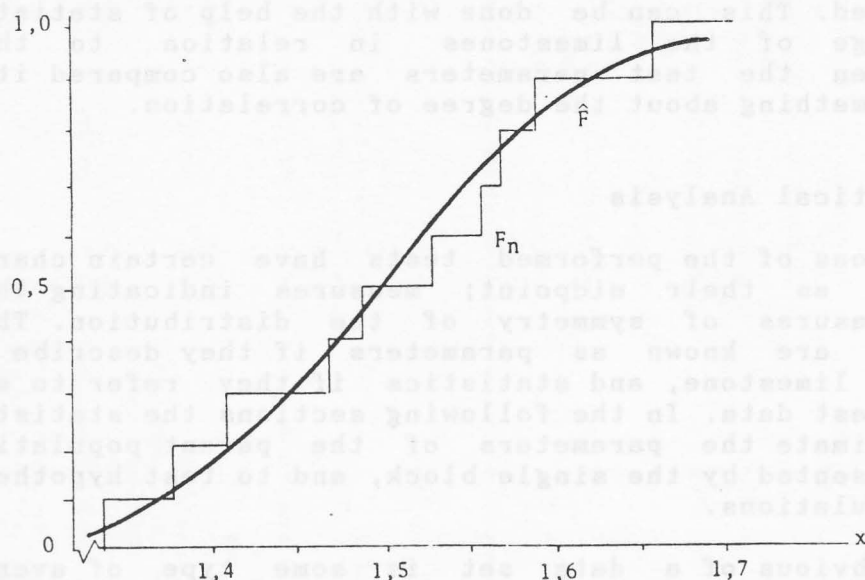


figure 5.1.1 the cumulative frequency distribution of a data set (xi) (van Soest 1983)

In case of a small number of data per set the most suitable test for normality is the one characterized as the empirical distribution function F_n (appendix C.1), which estimates the real distribution F . With this method it is possible to check the cumulative frequency distribution for normality with a certain unreliability (fig. 5.1.1). If the integrated quadratic difference between F_n and F is lower than a critical value, the distribution of the data set gives no cause for the population not to be normally distributed. This critical value is defined by the unreliability and the number of data in the set. If however the difference is higher than the normality of the data set is rejected and nothing can be said about the distribution of the population. In that event the median value is then taken as a representative average.

A measurement of the spread in the distributions is given by the calculation of the standard deviation for every set containing more than 2 data.

After application of the normality test it proved that 86 % of the data sets was normal distributed with a 10 % unreliability and 14 % of the data were not normal distributed. Hence in table 5.1.1 the average values are marked with an (*) are median values of negative tested data sets and those marked with a (+) are mean values of positive tested data sets. Behind them the averages, standard deviations and the number of data are given.

In case of the porosity of the Lower Oolite limestone (SL03) the difference between the three values was so high that it seemed not reasonable to define mean and deviation here.

5.2 Remarks on the average values

The UCS shows a wide range of values, which may be considered are representative for limestones with relative high moisture contents, in specific the weak limestones. An overview of all the stress vs. strain curves is given in figures 5.2.1-5.2.3, each plot containing 4-6 specimen of the same type. In figure 5.2.4 the relation of UCS vs. Geological age proves that not always the strength does increase with time. Four types (Permian limestone SL01, Silurian limestone SL05, Carboniferous limestone SL06 and Devonian limestone SL10) show deviating behaviour, which may be the result of external influences such as the relaxation of tectonic stresses or weathering processes.

The highest E-moduli are approximately 70 GPa and are therefore concurring to the expectation that pure calcite has the highest value, namely 85.0 GPa (Belikov 1967). The type of failure is for the stronger limestones generally an axial multi fracture, whereas for the weak ones mainly conical shaped end segments are formed along single shear planes. From figures 5.2.5

$\frac{1}{\sqrt{2}} \sqrt{\frac{1}{n} \sum_{i=1}^n (x_i - \bar{x})^2}$	$\frac{1}{\sqrt{2}} \sqrt{\frac{1}{n} \sum_{i=1}^n (x_i - \bar{x})^2}$	$\frac{1}{\sqrt{2}} \sqrt{\frac{1}{n} \sum_{i=1}^n (x_i - \bar{x})^2}$	$\frac{1}{\sqrt{2}} \sqrt{\frac{1}{n} \sum_{i=1}^n (x_i - \bar{x})^2}$
10000.0	9999.0	9998.0	9997.0
10001.0	9999.0	9998.0	9997.0
10002.0	9999.0	9998.0	9997.0
10003.0	9999.0	9998.0	9997.0
10004.0	9999.0	9998.0	9997.0
10005.0	9999.0	9998.0	9997.0
10006.0	9999.0	9998.0	9997.0
10007.0	9999.0	9998.0	9997.0
10008.0	9999.0	9998.0	9997.0
10009.0	9999.0	9998.0	9997.0
10010.0	9999.0	9998.0	9997.0
10011.0	9999.0	9998.0	9997.0
10012.0	9999.0	9998.0	9997.0
10013.0	9999.0	9998.0	9997.0
10014.0	9999.0	9998.0	9997.0
10015.0	9999.0	9998.0	9997.0
10016.0	9999.0	9998.0	9997.0
10017.0	9999.0	9998.0	9997.0
10018.0	9999.0	9998.0	9997.0
10019.0	9999.0	9998.0	9997.0
10020.0	9999.0	9998.0	9997.0

In case of a small number of data per set, the most suitable test for normality is the one characterized as the empirical distribution function $F_n(x)$ which estimates the real distribution $F(x)$. With this method it is possible to check the cumulative frequency distribution for normality with a certain unreliability (fig. 5.1.1). If the interested quadratic difference between F_n and F is lower than a critical value, the distribution of the data set gives no cause for the population not to be normally distributed. This critical value is defined by the unreliability and the number of data in the set. If however the difference is higher than the normality of the data set is rejected and nothing can be said about the distribution of the population. In that event the median value is then taken as a representative average.

A measurement of the spread in the distributions is given by the calculation of the standard deviation for every set containing more than 2 data.

After application of the normality test it proved that 88 % of the data sets were normal distributed with a 10 % unreliability and 12 % of the data were not normal distributed. Hence in table 5.1.1 the average values are marked with an (*) are median values of negative tested data sets and those marked with a (•) are mean values of positive tested data sets. Behind them the average standard deviation and the number of data are given.

In case of the porosity of the lower Oolite limestone (E102) the difference between the three values was so high that it seemed not reasonable to define mean and deviation here.

5.2 Remarks on the average values

The UCS shows a wide range of values, which may be considered as representative for limestones with relative high moisture contents. In specific the weak limestones. An overview of all the stress vs. strain curves is given in figure 5.2.1. 5.2.2 each plot containing 4-8 specimens of the same type. In figure 5.2.4 the relation of UCS vs. Geological age proves that not always the strength does increase with time. Four types (Permian limestone E101, Silurian limestone E108, Carboniferous limestone E109 and Devonian limestone E110) show deviating behaviour, which may be the result of external influences such as the relaxation of tectonic stresses or weathering processes.

The highest E-moduli are approximately 70 GPa and are therefore concerning to the expectation that pure calcite has the highest value, namely 85.6 GPa (Belikov 1987). The type of failure is for the strongest limestones generally an axial split fracture, whereas for the weak ones mainly conical shaped and segments are formed along single shear planes. From figure 5.2.3

- 5.2.7 to typical E-moduli are viewed for all the samples.

The Poisson ratio's vary from 0.23 to 0.34 which is in accordance with the mentioned limestone ranges in the literature (Lama & Vutukuri 1978).

The ultra sonic velocities are can be related with the velocity of calcite. The quality of the limestone can be indexed by means of a ratio (Fourmaintraux 1976) :

$$IQ \% = \frac{V_{\text{sample}}}{V_{\text{calcite}}} * 100 \% \quad \text{and } V_{\text{calcite}} = 6600 \text{ m/s}$$

Experiments have established that the IQ is affected by pores and fissures. Typical values for strong limestones range from 6000 to 6500 m/s .

The brazilian tensile strength shows only for the "Maassteen" (SL07) an unexpected low value. The reason is unknown. From the stress vs. strain plots in fig. 5.2.8 - 5.2.10 it is seen that the stability of the curve decreases with increasing strain. The reason is the inaccuracy of the load transducer. The weak Limburgian Marl shows an enormous frequency, which is due to the large amount of strain measurements at low stress.

The measured porosity is an effective porosity, hence permeable limestones, such as the Limburgian Marl (SL12) and the Magnesian Limestone (SL01), will approach the total porosities to a higher degree than the nearly impermeable ones.

The zero permeability values of the stronger limestones indicates that nothing could be measured. The general permeabilities of these limestones should be, according to the literature (Brace 1978) between 10^{-5} and 10^{-13} m/s.

The slake durability can be rated with the classification of Gamble (1971) Concerning the data the lowest durability is then rated as "medium durability" (table 5.2.1).

groupname	% retained after two cycles
very high durability	> 98
high durability	95-98
medium high durability	85-95
medium durability	60-85
low durability	30-60
very low durability	< 30

table 5.2.1 Gamble's slake durability classification

TIME-STRENGTH RELATION

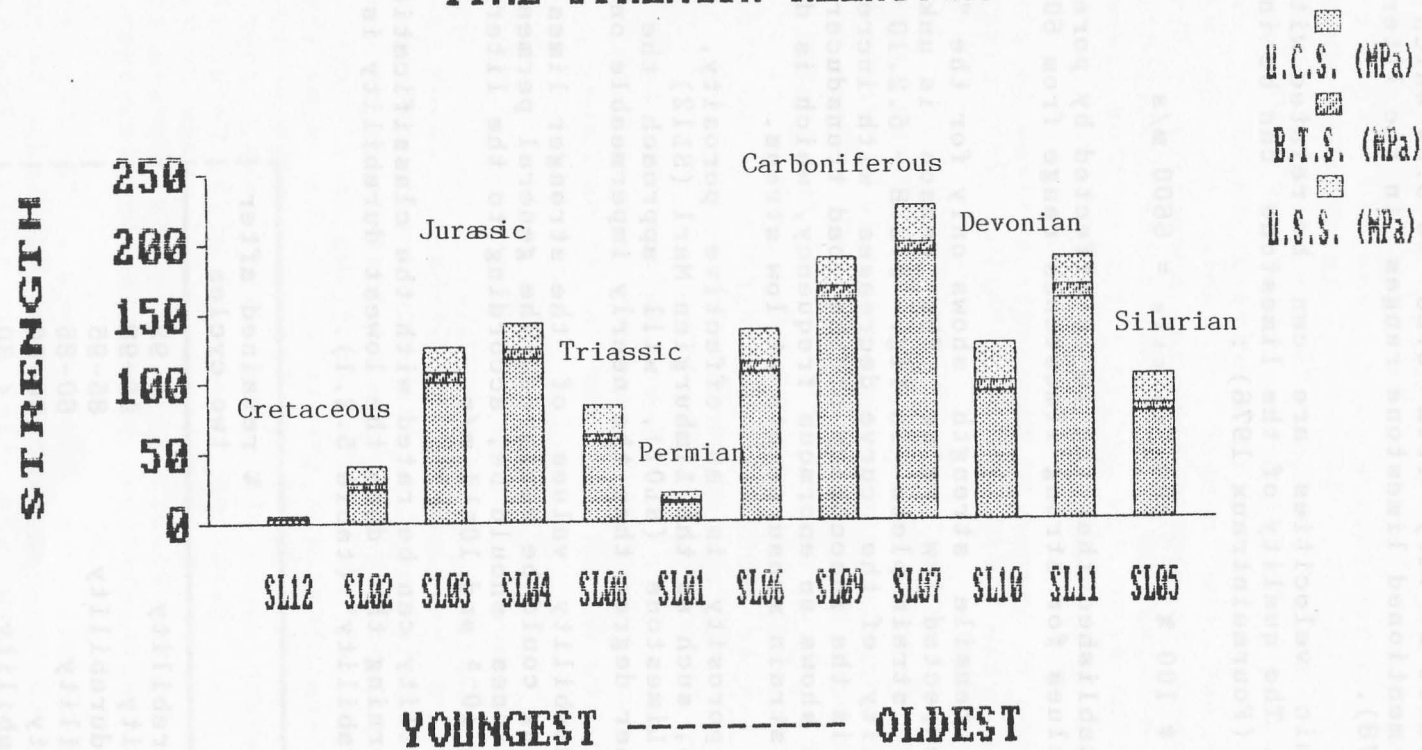


figure 5.2.4 The time-strength relation for 12 types of limestone with increasing age from left to right

The dry densities are all lower than the absolute density of pure calcium carbonate; specific gravity of CaCO_3 is ~ 2.71 (Handbook of Chemistry and Physics 1982). The differences in dry density are in the first place explained by pores and fissures and secondly by the contaminant minerals of the samples, with different specific gravities; MgCO_3 (magnesium) has a specific gravity of ~ 2.95 . This causes the gravity of the Magnesian Limestone (SL02) to be slightly higher than normal.

From the X-ray diffraction it is apparent that not all the sampled limestones are indeed pure limestones; the Magnesian Limestone (SL01), the Upper Oolite Limestone (SL04) and the Muschelkalk (SL08) have a CaCO_3 content between 50 % and 90 %, and thus actually have to be named "impure limestones" following the geological classification of Leighton & Pendexter.

The lowest value of the CR-Index is estimated to be in the range of 65-70 (fig. 5.3.2). Deviating values of the Carbonate Resistance Index are the Upper Chalk (SL02) and the "Maassteen" (SL07). It is found from the results that the solubility is related to the effective porosity of the rock.

5.3 Correlation of Parameters

When the parameters are mutually compared two types of functions are suggested from the correlation of the plots; linear and logarithmic functions. If the correlation coefficient for such a function is greater than 0.8 it is assumed that the parameters are related. In the following sections correlations are made between the most important parameters.

Generally there are three groups distinguished by the sort of parameters compared; the mechanical parameters, like UCS, BTS, E-modulus and the physical parameters like porosity, density, sonic velocity and moisture content.

In the first group the mechanical parameters are compared with each other, and in figure 5.3.1 linear correlations are evident; If the UCS increases the BTS and E-modulus increase proportionally.

In the second group mechanical parameters are correlated with physical ones (fig. 5.3.2). It shows that these relations have a strong tendency towards logarithmic functions, because their correlation coefficient may be as high as 0.96. The logarithmic functions have asymptotes at X and Y values. For example the sensitivity of UCS is the highest for very low porosity values and therefore the relation has a vertical asymptote. It is confirmed in the literature (Dearman 1982) that this is true (fig. 5.3.3) for dolerites.

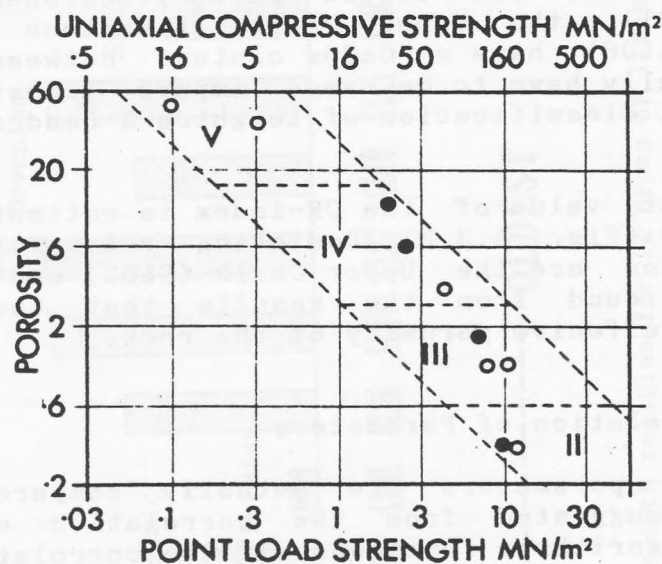


Fig.1 Relationship between strength and porosity in weathered dolerites

figure 5.3.3 The relation between Porosity and UCS (Dearman 1982)

The third group regards physical vs. physical parameters, such as dry density vs. porosity (fig. 5.3.4). For this example a linear correlation is almost absolute. When reviewing the literature it was found that this is correct (Goodman 1980) for the relation porosity vs. dry density :

$$\gamma_{dry} = \gamma_w \cdot G \cdot (1-n)$$

were γ_{dry} = dry density
 γ_w = wet density
 G = specific gravity
 n = porosity

The other correlations in this group exhibit the same linearity.

Finally the correlation of grain sizes with mechanical and physical properties did not exist for the sample data. Correlation coefficients were between -0.5 and 0.5 .

The third group, regarding physical vs. physical parameters, such as dry density vs. porosity (Fig. 5.2.4), for this example a linear correlation is almost absolute. When reviewing the literature it was found that this is correct (Goodman 1962) for the relation porosity vs. dry density:

$$Y_{dr} = Y_w - G(1-p) \quad \text{were } Y_{dr} = \text{dry density}$$

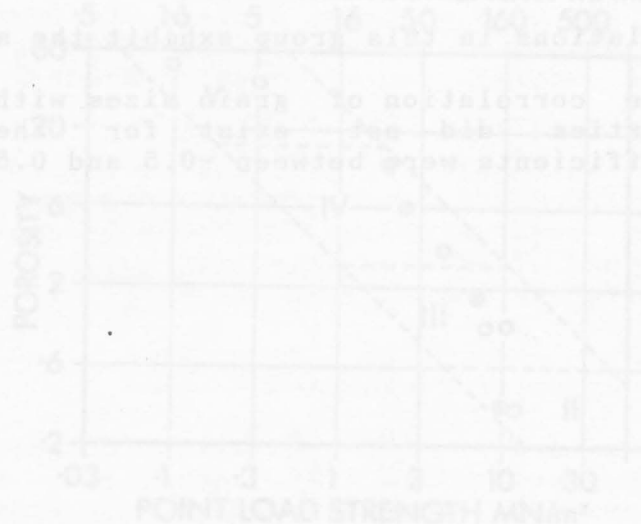
$$Y_w = \text{wet density}$$

$$G = \text{specific gravity}$$

$$p = \text{porosity}$$

UNIAXIAL COMPRESSIVE STRENGTH MINN

The other correlation in this group exhibits the same linearity. Finally the correlation of grain sizes with mechanical and physical properties did not exist for the sample data. Correlation coefficients were between -0.5 and 0.5.



- Dolomite, New England Quarry, S. Devon
- Contact metamorphosed dolomite, Maiden Quarry, S. Devon, England
- Weathering grade

Fig. 5.2.3 Relationship between strength and porosity in weathered dolomite

Figure 5.2.3 The relation between porosity and uniaxial compressive strength (Goodman 1962)

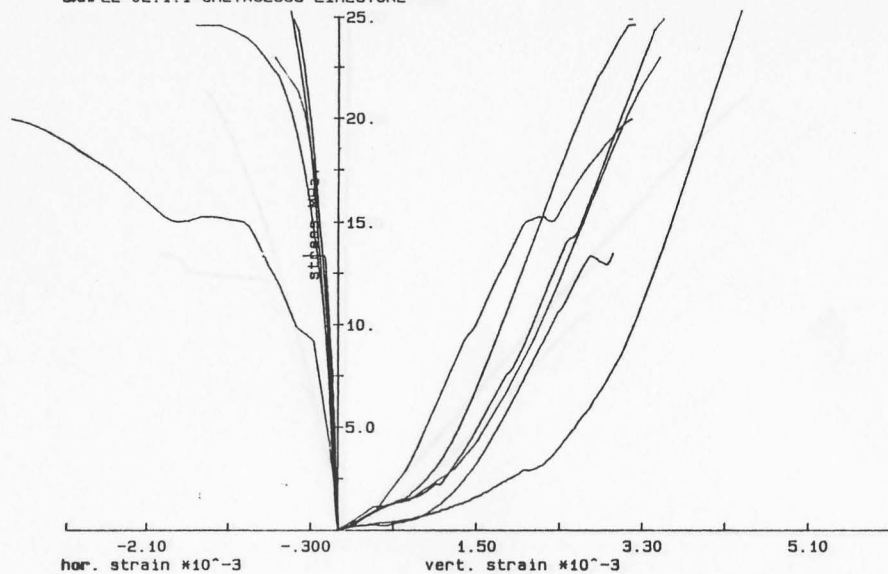
CODE	U.C.S.	(MPa)		E-MODULUS	(GPa)		POISSON RATIO		SONIC VEL.	(M/S)	
SAMPLE	Avq.	[S.D.]	Nx	Avq.	[S.D.]	Nx	Avq.	[S.D.]	Avq.	[S.D.]	Nx
SL01	12.82 +	[1.75]	6	4.48 *	[.91]	6	.31 *	[.20]	2580 +	[110]	6
SL02	24.76 *	[4.97]	6	11.15 +	[1.92]	6	.23 +	[.11]	4198 +	[117]	6
SL03	100.46 +	[12.56]	6	40.15 +	[1.42]	6	.31 +	[.02]	5560 +	[26]	6
SL04	116.98	[21.73]	5	32.45	[2.96]	5	.31	[.03]	5169 +	[123]	6
SL05	75.51 +	[3.63]	6	63.13 +	[.85]	6	.34 +	[.02]	5535 *	[184]	6
SL06	105.40	[48.40]	4	63.72	[1.41]	4	.32	[.02]	6358 +	[77]	6
SL07	190.43 +	[4.46]	6	70.34 +	[.34]	6	.31 +	[.01]	6553 +	[18]	6
SL08	57.43 +	[17.99]	6	14.27 +	[2.20]	6	.35 *	[.19]	4137 *	[814]	6
SL09	157.08 +	[5.12]	6	69.92 +	[.69]	6	.33 +	[.02]	6440 +	[18]	6
SL10	91.31 *	[17.30]	6	55.12 +	[3.04]	6	.29 +	[.05]	6139 +	[75]	6
SL11	157.48 +	[12.21]	6	67.23 +	[5.28]	6	.32 +	[.02]	6183 +	[48]	6
SL12	3.52 +	[.41]	6	1.21 +	[.11]	6	.25 +	[.04]	1766 +	[22]	6

CODE	B.T.S.	(MPa)		U.S.S.	(MPa)		POROSITY	(%)		PERMEAB.	(mDarcy)	SLAKE DURABILITY	
SAMPLE	Avq.	[S.D.]	Nx	Avq.	[S.D.]	Nx	Avq.	[S.D.]	Nx	Avq.	[S.D.]	Nx	INDEX
SL01	2.17 *	[.55]	6	6.14 +	[1.6]	6	37.97	[.18]	3	2.88	[.60]	3	80.91
SL02	2.95 +	[.83]	6	13.16 +	[2.4]	6	15.70	[.82]	3	0		3	98.03
SL03	6.24 +	[.47]	6	20.08 +	[4.3]	6				0		3	98.78
SL04	6.88 +	[1.00]	6	19.64 +	[2.5]	6	2.00	[.18]	3	0		3	99.65
SL05	6.18 +	[.46]	6	20.94 *	[5.4]	6	1.08	[.09]	3	0		3	98.86
SL06	7.75 +	[1.80]	6	24.04 +	[3.5]	6	.57	[.07]	3	0		3	98.11
SL07	7.34 +	[.65]	6	25.97	[3.9]	5	.68	[.18]	3	0		3	99.32
SL08	5.79 +	[1.09]	6	19.66 *	[2.7]	6	5.42	[1.21]	3	0		3	96.31
SL09	9.13 *	[1.65]	6	21.63 +	[1.2]	6	.97	[.08]	3	0		3	99.38
SL10	6.31 +	[.83]	6	28.53 +	[4.3]	6	.59	[.08]	3	0		3	98.99
SL11	9.09 +	[.97]	6	21.11 +	[2.1]	6	.79	[.04]	3	0		3	99.48
SL12	.38 +	[.03]	6	.83 +	[.2]	6	50.05	[.09]	3	5450		2	71.41

CODE	MOIST.CONT	(%)		WET DENSITY	(TON/M3)		DRY DENSITY	(TON/M3)		CaCO3	MgCO3	CR-INDEX
SAMPLE	Avq.	[S.D.]	Nx	Avq.	[S.D.]	Nx	Avq.	[S.D.]	Nx	(%)	(%)	Avq. of 2
SL01	17.04	[.31]	3	2.12	[.01]	3	1.76	[.01]	3	62.18	37.92	158.7
SL02	5.15	[.29]	3	2.40	[.02]	3	2.28	[.02]	3	97.60	.90	72.8
SL03	1.48	[.12]	3	2.62	[.01]	3	2.58	[.02]	3	92.66	1.19	85.8
SL04	1.94	[.25]	3	2.58	[.01]	3	2.53	[.02]	3	83.64	.99	78.2
SL05	.03	[.01]	3	2.68	[.00]	3	2.68	[.00]	3	97.41	1.10	80.7
SL06	.18	[.04]	3	2.68	[.00]	3	2.67	[.00]	3	98.40	1.13	76.9
SL07	.77	[.02]	3	2.69	[.01]	3	2.68	[.01]	3	99.37	.74	90.0
SL08	.70	[.08]	3	2.57	[.01]	3	2.55	[.01]	3	82.01	4.29	111.1
SL09	.03	[.02]	3	2.67	[.01]	3	2.66	[.01]	3	97.13	1.51	78.5
SL10	.09	[.04]	3	2.69	[.00]	3	2.69	[.00]	3	91.00	2.00	77.2
SL11	.17	[.02]	3	2.69	[.00]	3	2.68	[.00]	3	96.71	1.99	76.4
SL12	22.05	[.13]	3	1.71	[.01]	3	1.34	[.00]	3	96.64	1.70	262.3

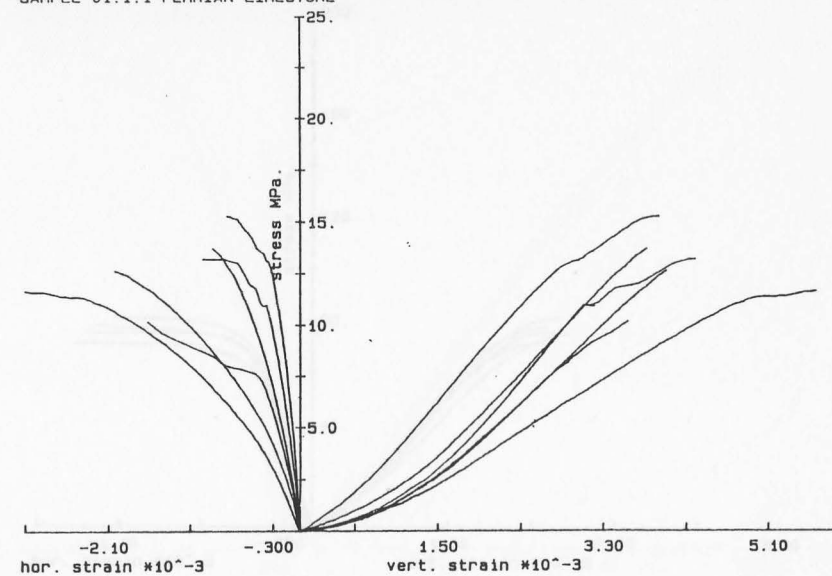
table 5.1.1

SAMPLE 02.1.1 CRETACEOUS LIMESTONE



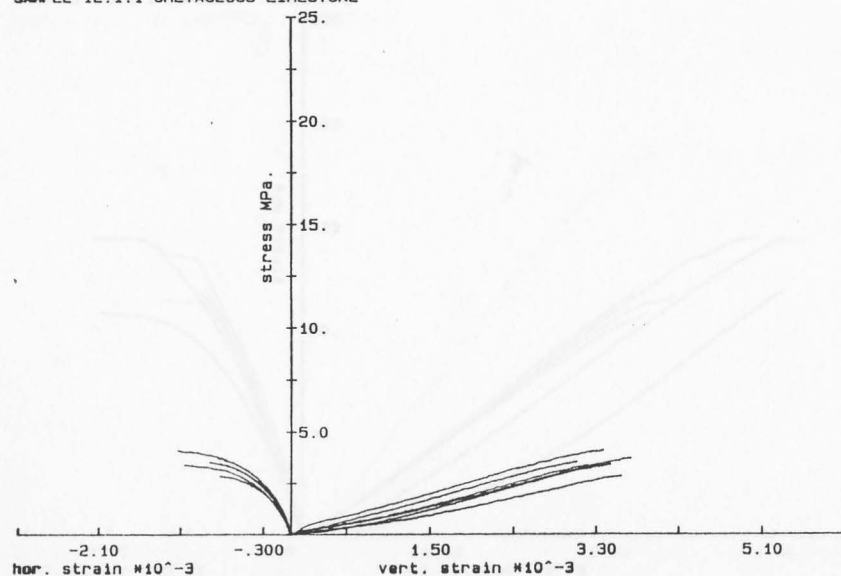
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SAMPLE 01.1.1 PERMIAN LIMESTONE



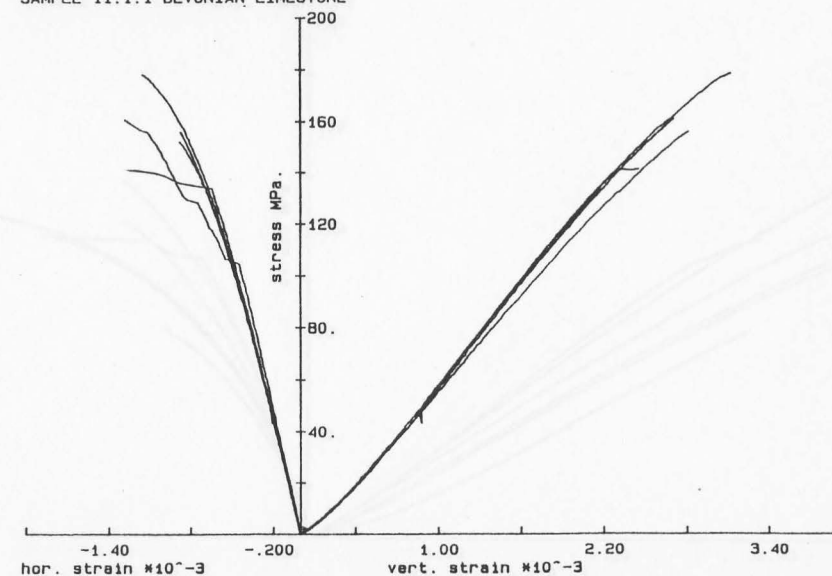
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SAMPLE 12.1.1 CRETACEOUS LIMESTONE



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SAMPLE 11.1.1 DEVONIAN LIMESTONE



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Fig. 1. Dependence of the rate of polymerization on the concentration of the initiator.



Fig. 2. Dependence of the rate of polymerization on the concentration of the monomer.



Fig. 3. Dependence of the rate of polymerization on the concentration of the initiator.

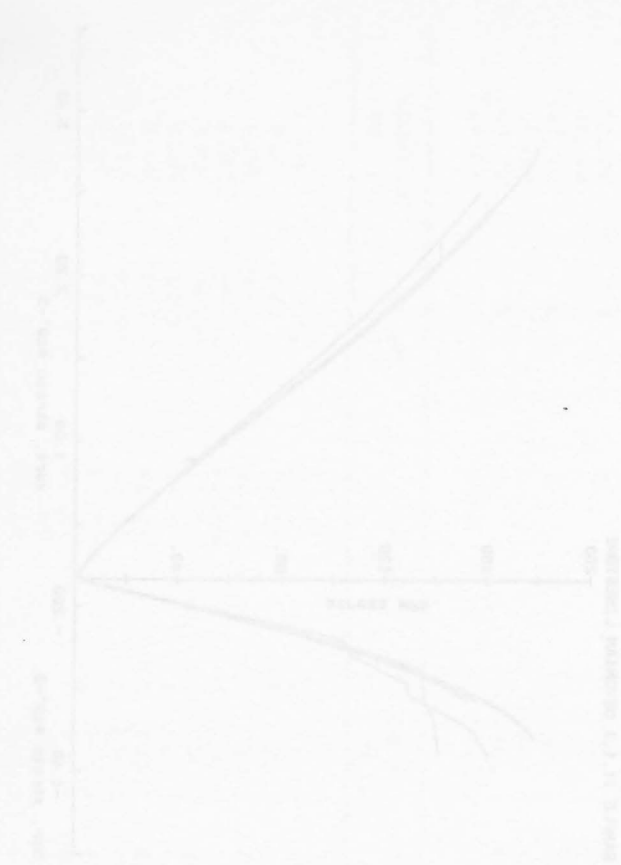
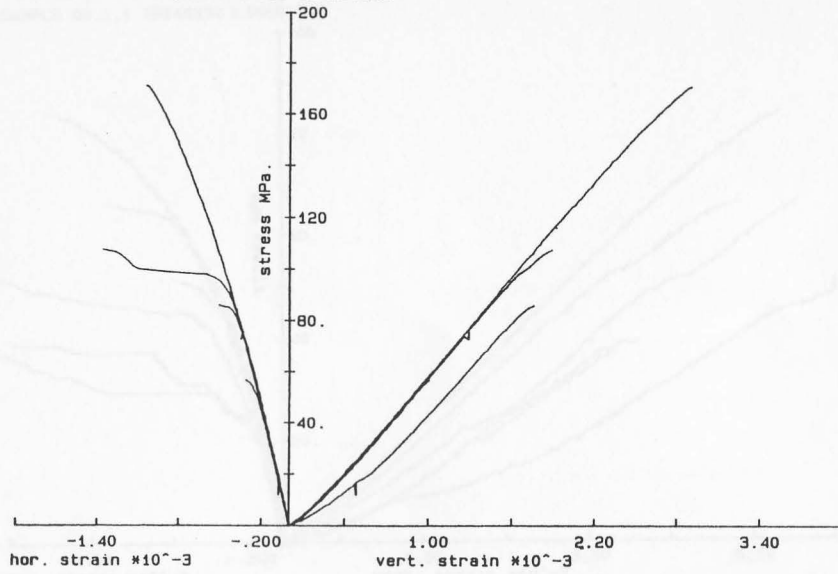


Fig. 4. Dependence of the rate of polymerization on the concentration of the monomer.

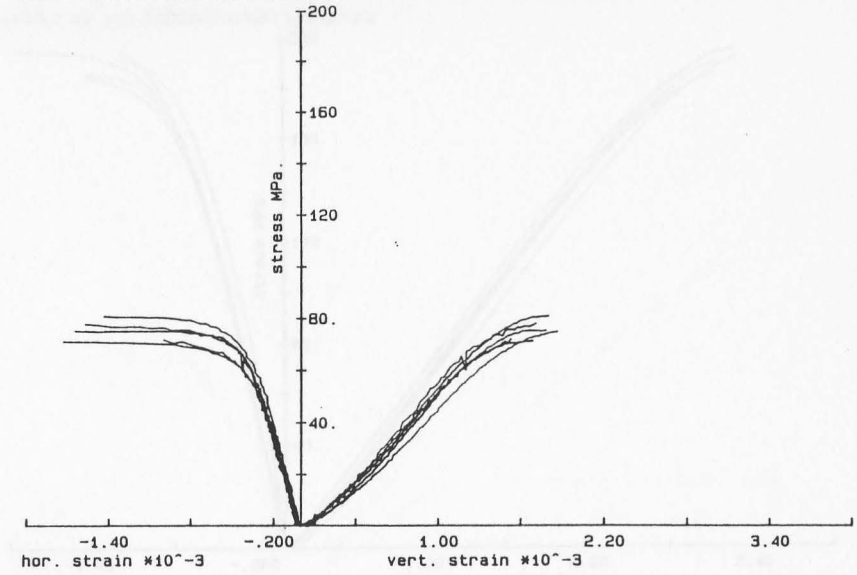


SAMPLE 06.1.2 CARBONIFEROUS LIMESTONE



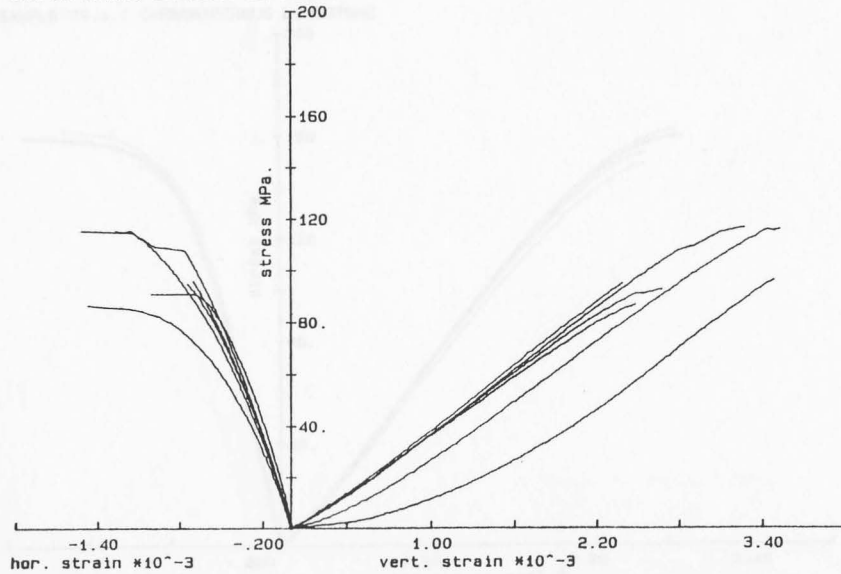
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SAMPLE 05.1.1 SILURIAN LIMESTONE



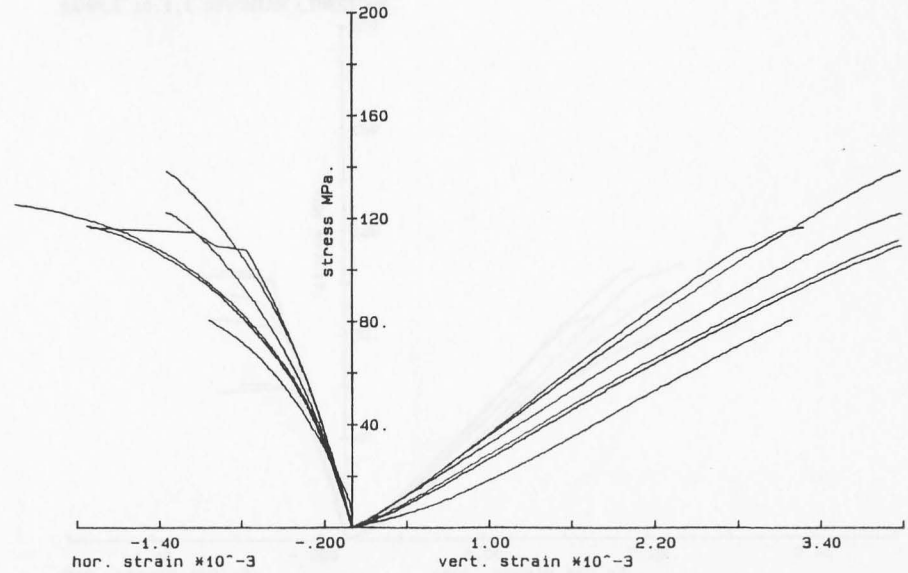
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SAMPLE 03.1.1 JURASSIC LIMESTONE



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SAMPLE 04.1.1 JURASSIC LIMESTONE



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Figure 2.5.1. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.

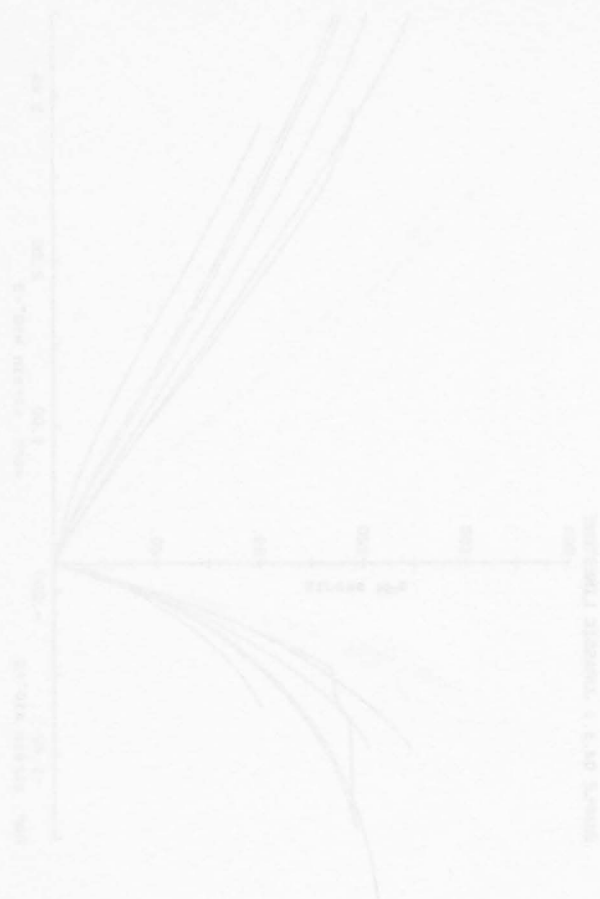


Figure 2.5.2. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.

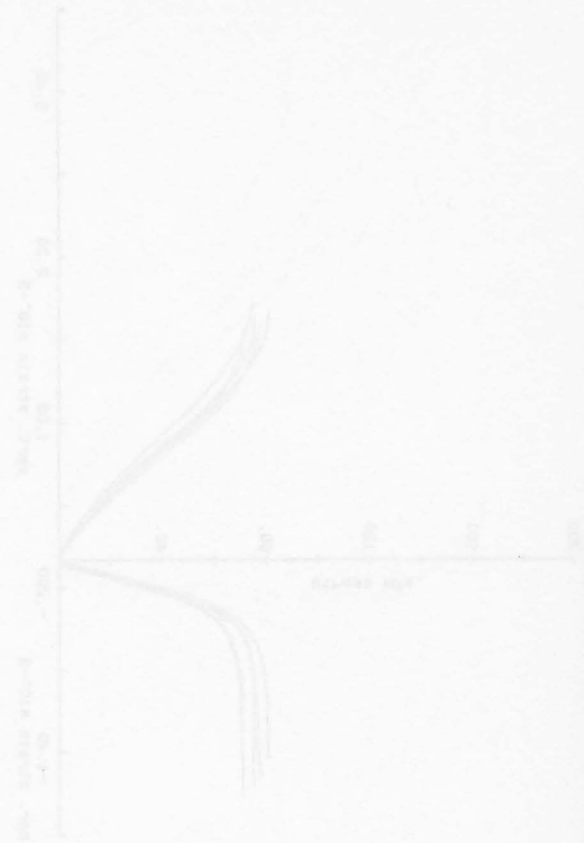
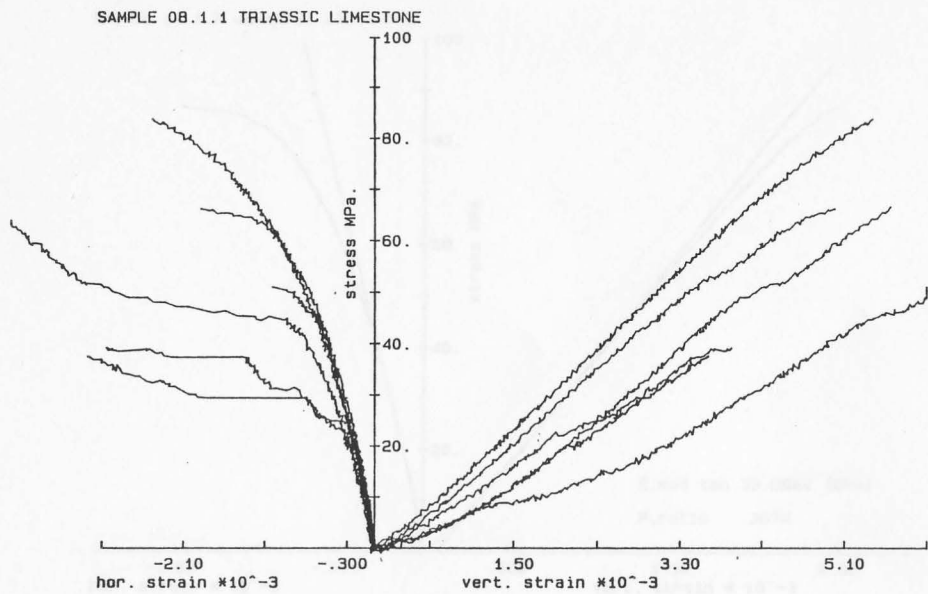


Figure 2.5.3. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.

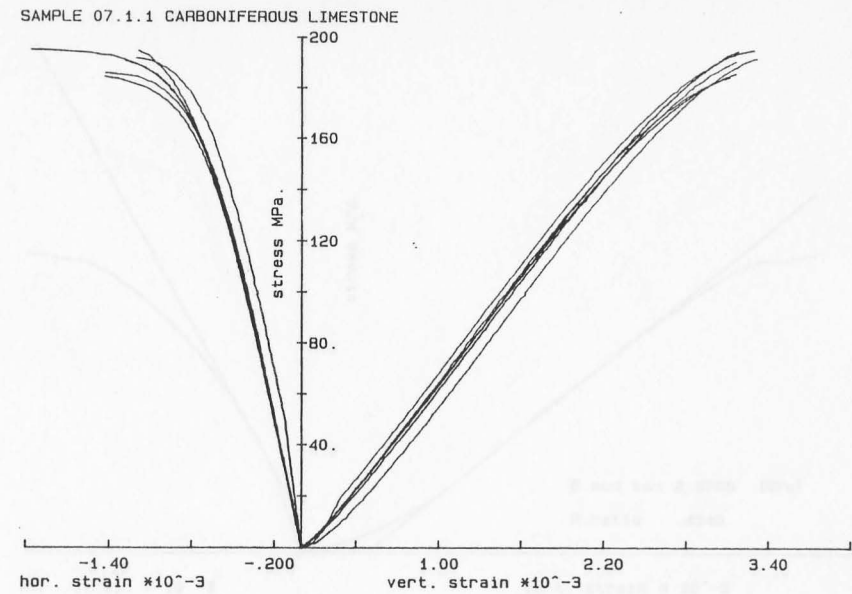


Figure 2.5.4. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.

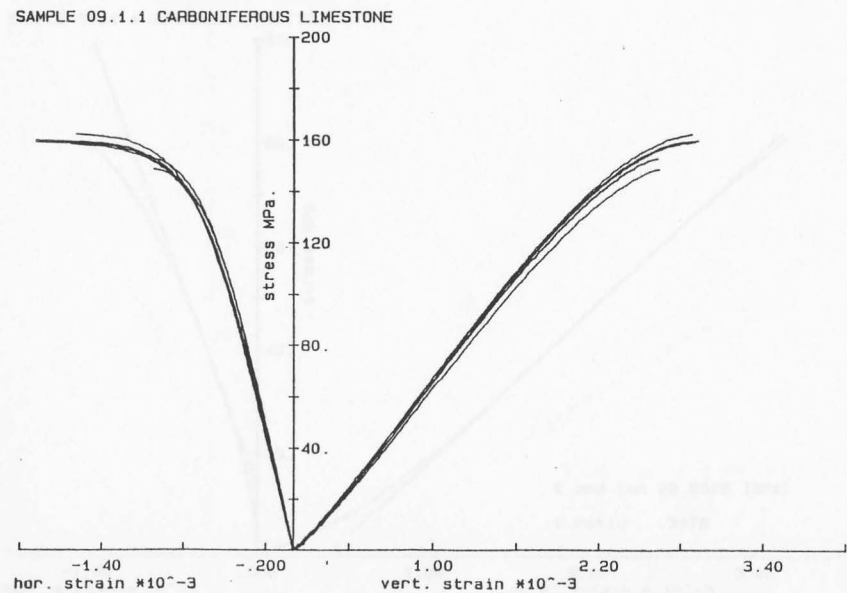




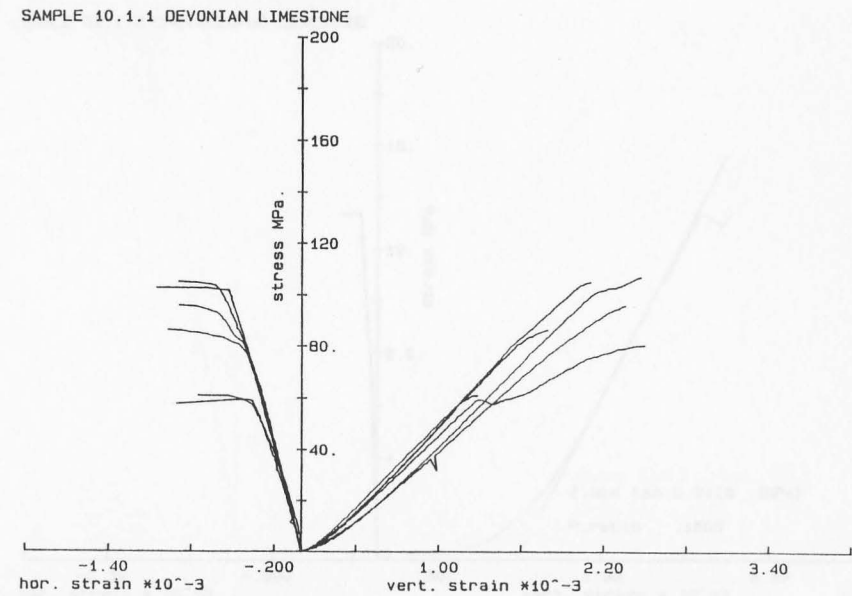
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Fig. 1. Dependence of the critical temperature on the concentration of the solution.

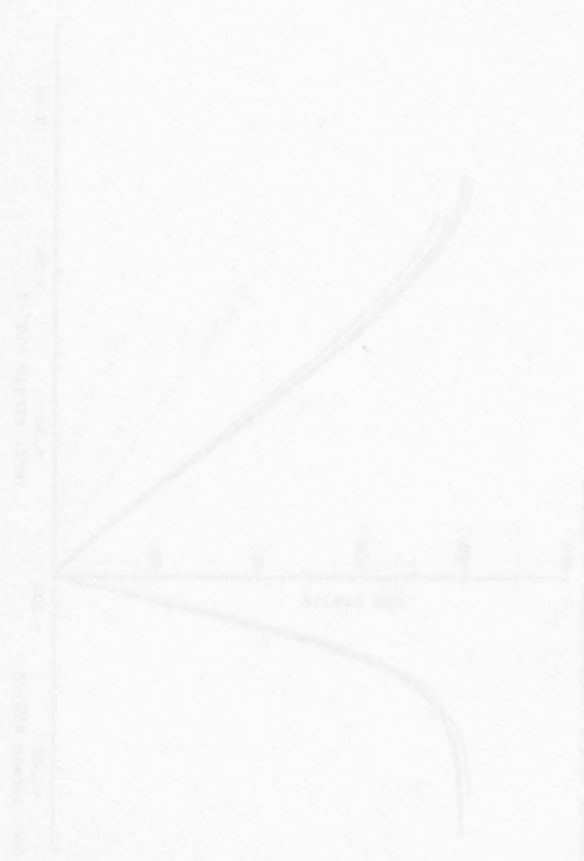


Fig. 2. Dependence of the critical temperature on the concentration of the solution.



Fig. 3. Dependence of the critical temperature on the concentration of the solution.

Fig. 4. Dependence of the critical temperature on the concentration of the solution.

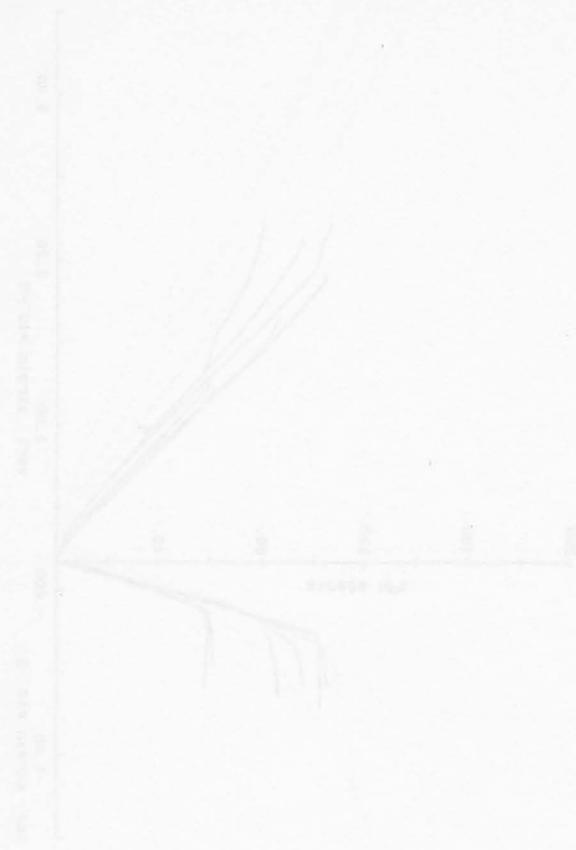


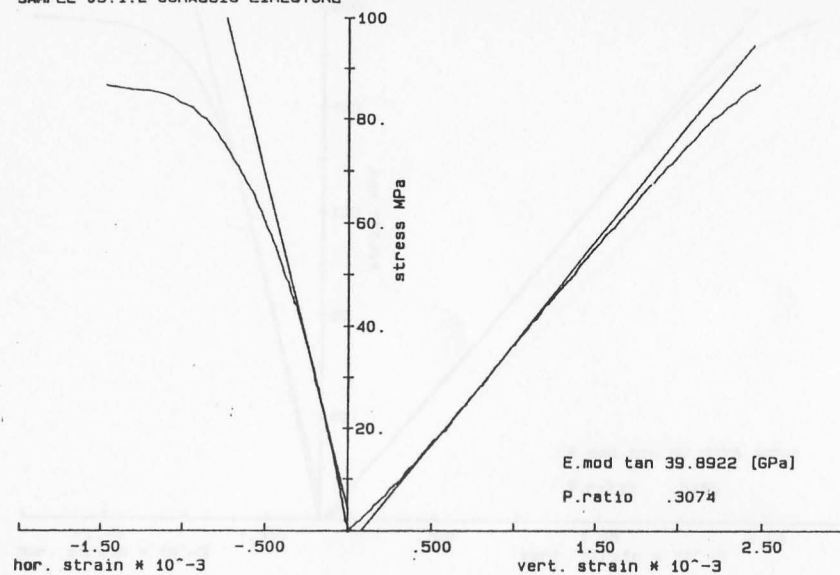
Fig. 5. Dependence of the critical temperature on the concentration of the solution.

Fig. 6. Dependence of the critical temperature on the concentration of the solution.



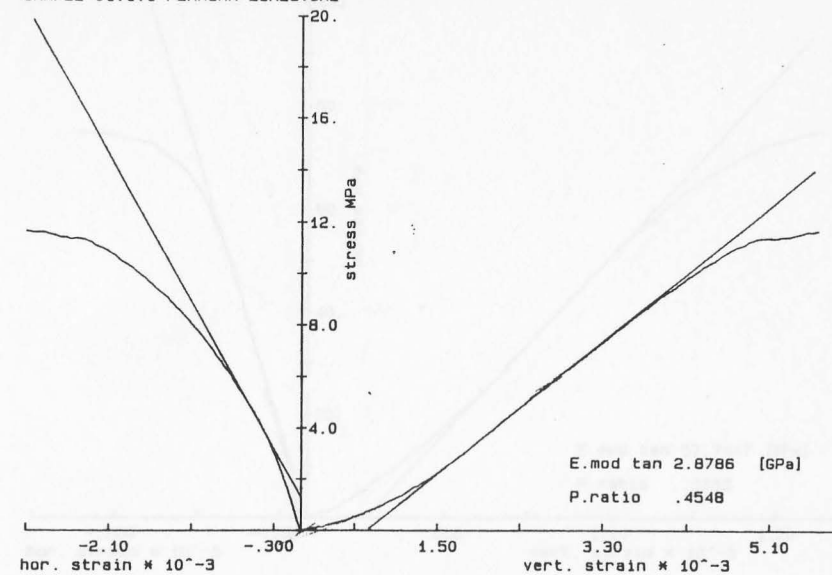
Fig. 7. Dependence of the critical temperature on the concentration of the solution.

SAMPLE 03.1.2 JURASSIC LIMESTONE



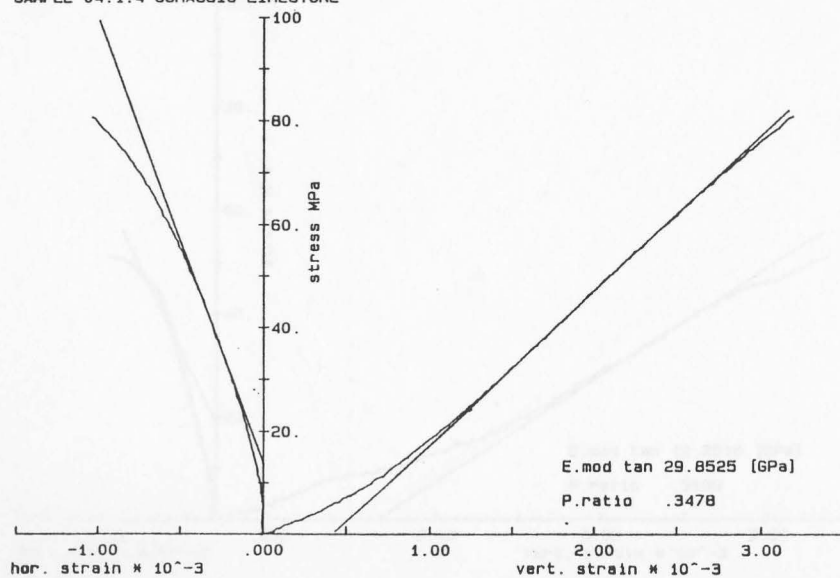
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SAMPLE 01.1.1 PERMIAN LIMESTONE



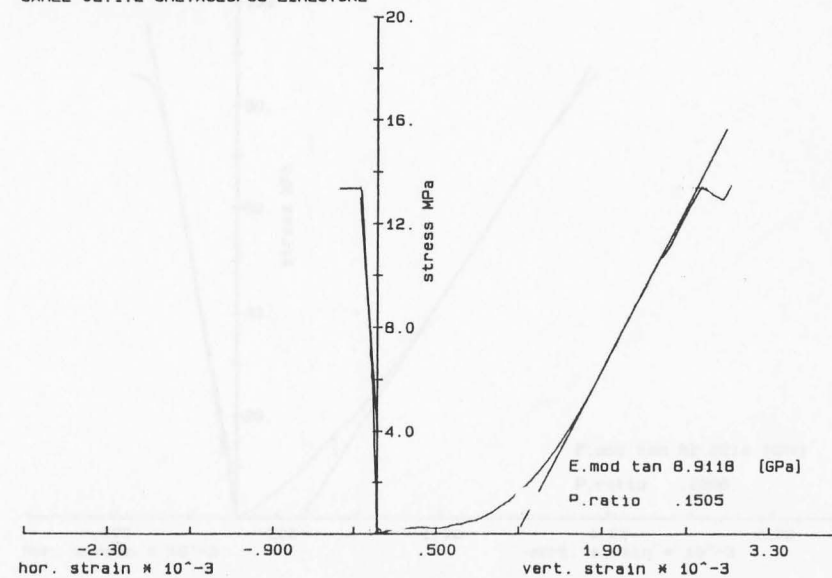
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SAMPLE 04.1.4 JURASSIC LIMESTONE



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SAMPLE 02.1.2 CRETACEOUS LIMESTONE



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Figure 2.4.1: Plot of $\ln(\frac{1}{1-x})$ versus x for x from 0 to 1. The curve starts at (0,0) and increases monotonically, passing through (0.5, 0.693).



Figure 2.4.2: Plot of $\ln(\frac{1}{1-x})$ versus x for x from 0 to 1. The curve starts at (0,0) and increases monotonically, passing through (0.5, 0.693).

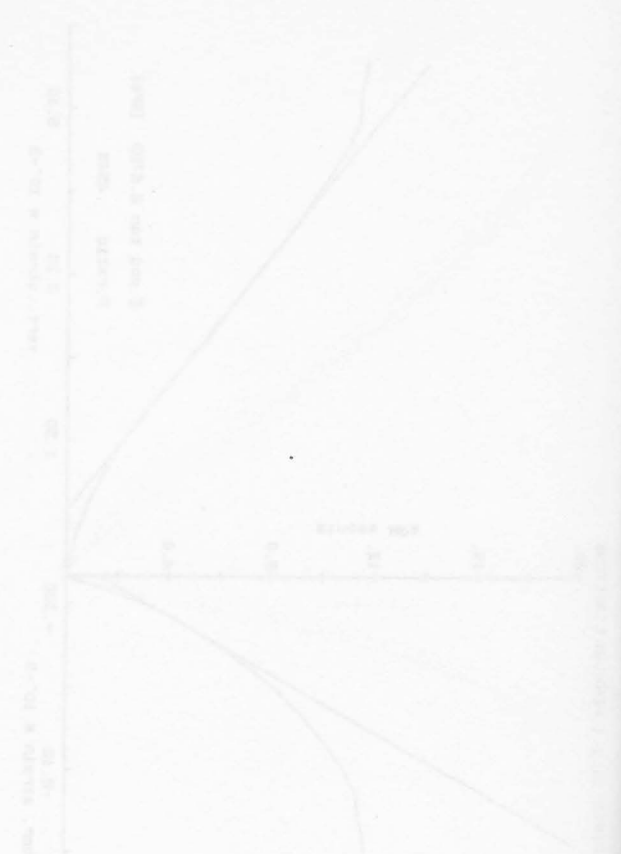
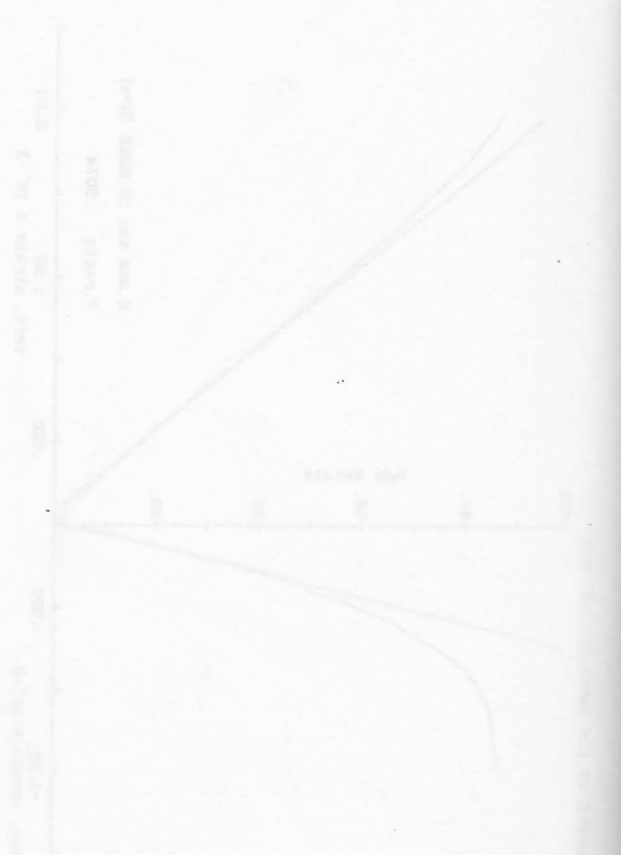
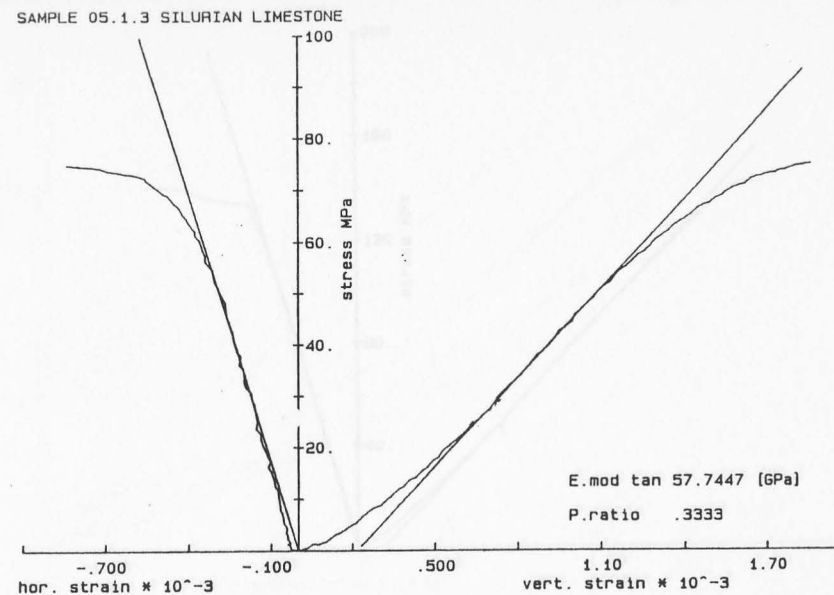
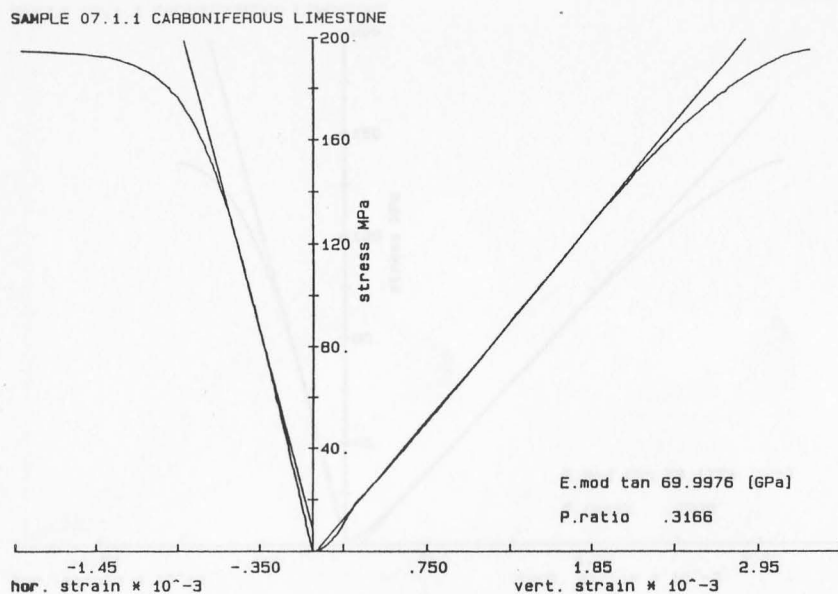


Figure 2.4.3: Plot of $\ln(\frac{1}{1-x})$ versus x for x from 0 to 1. The curve starts at (0,0) and increases monotonically, passing through (0.5, 0.693).



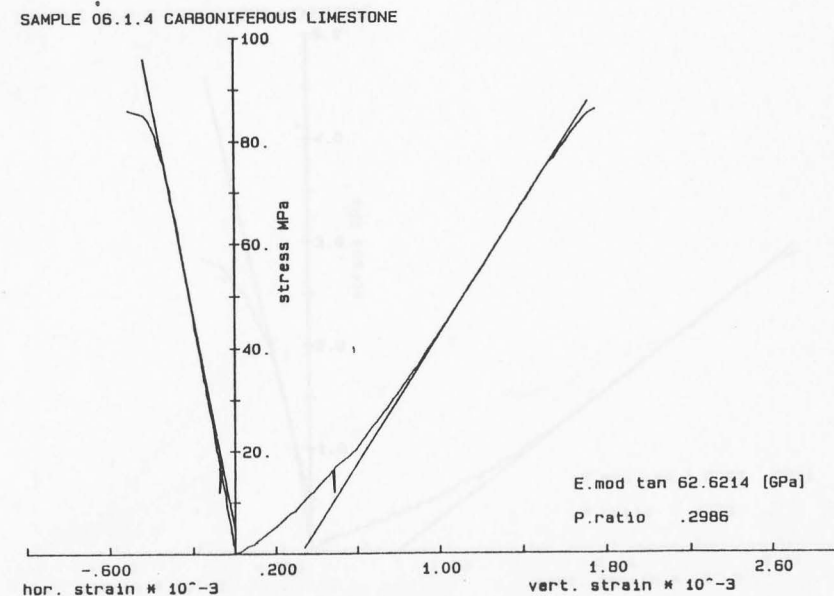
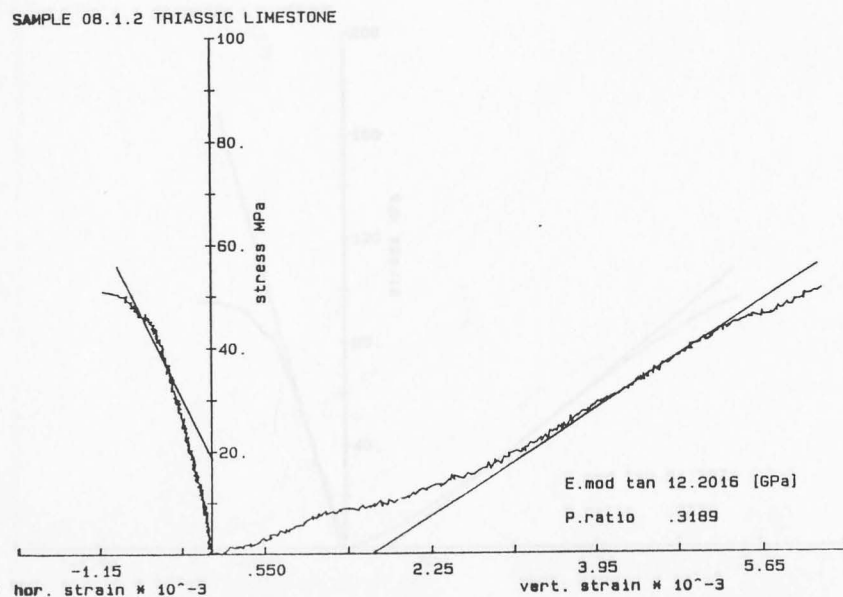
Figure 2.4.4: Plot of $\ln(\frac{1}{1-x})$ versus x for x from 0 to 1. The curve starts at (0,0) and increases monotonically, passing through (0.5, 0.693).





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Figure 2.3.2. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.



Figure 2.3.3. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.

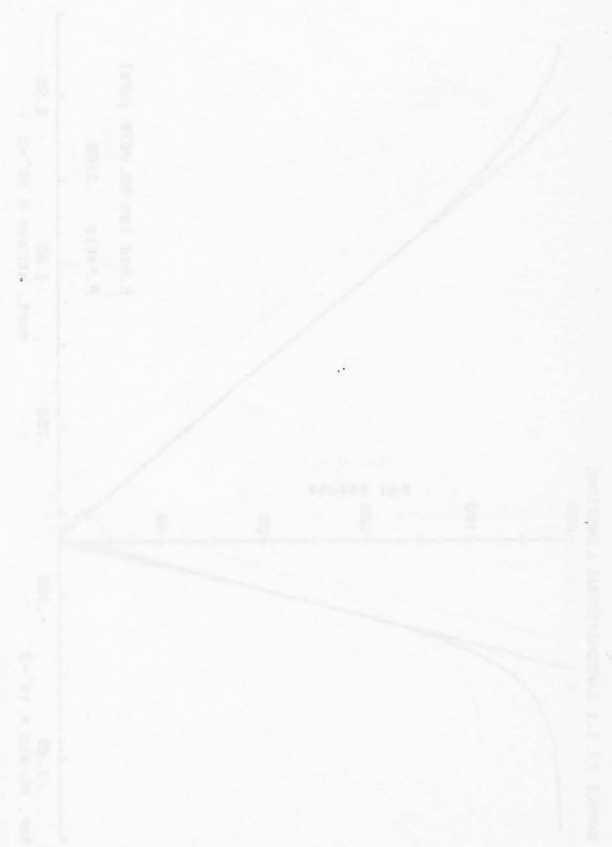


Figure 2.3.4. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.

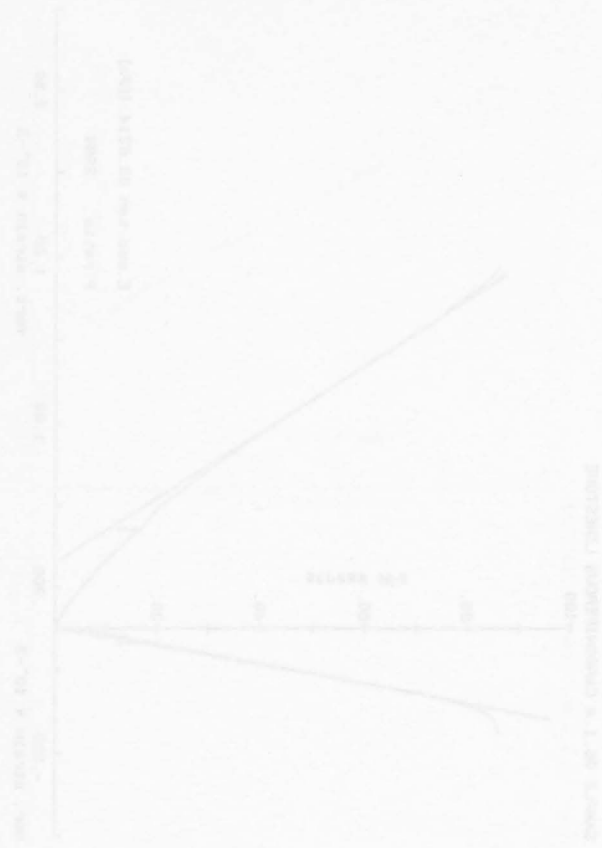
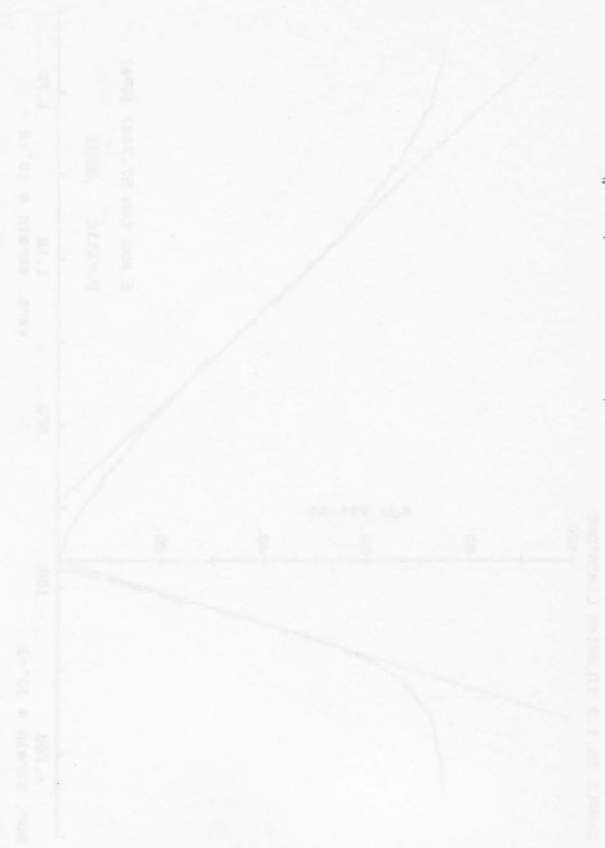
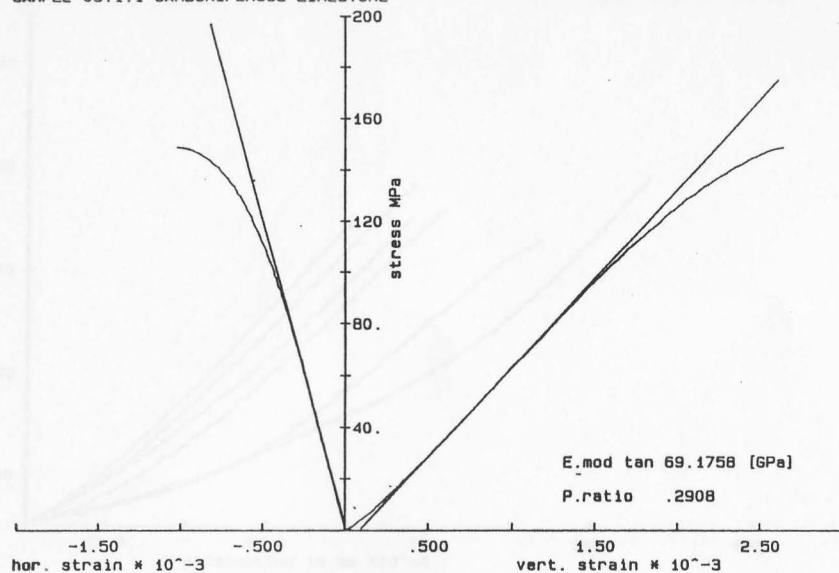


Figure 2.3.5. Polymerization of styrene initiated by azobisisobutyronitrile (AIBN) in benzene at 60°C.

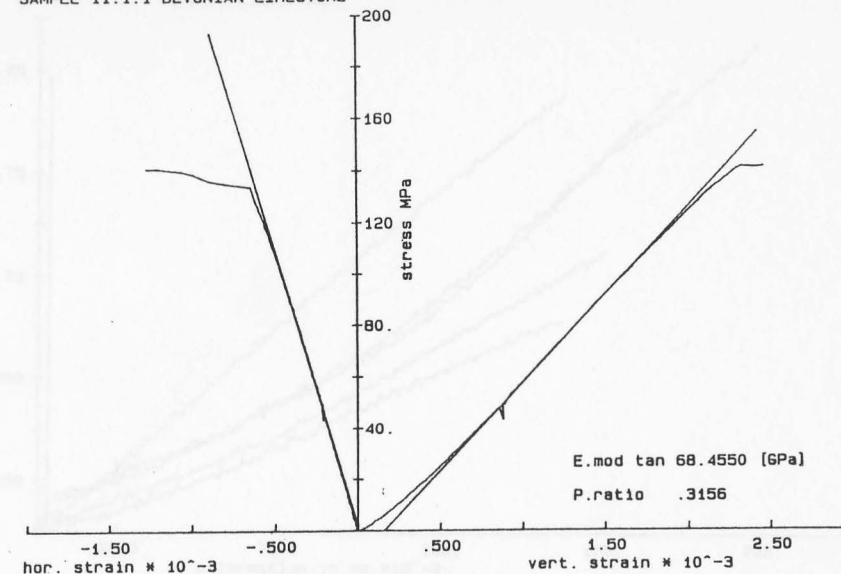


SAMPLE 09.1.1 CARBONIFEROUS LIMESTONE



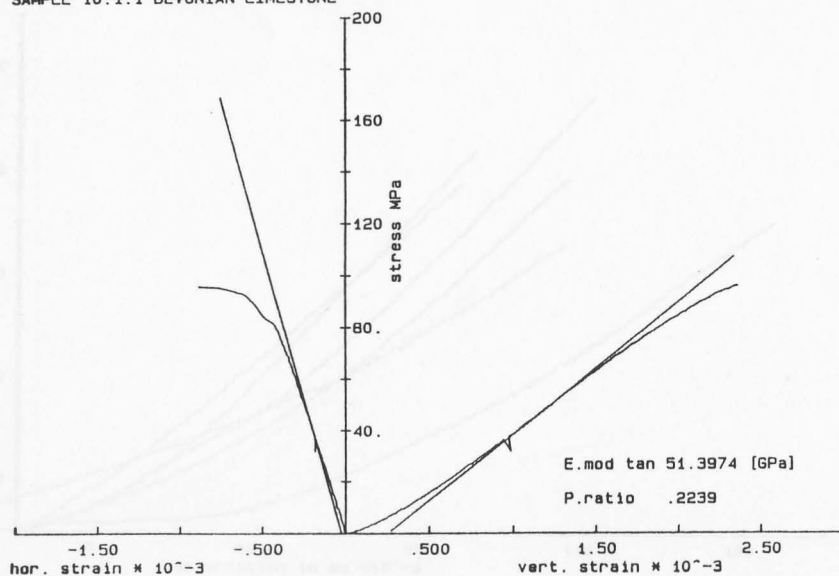
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SAMPLE 11.1.1 DEVONIAN LIMESTONE



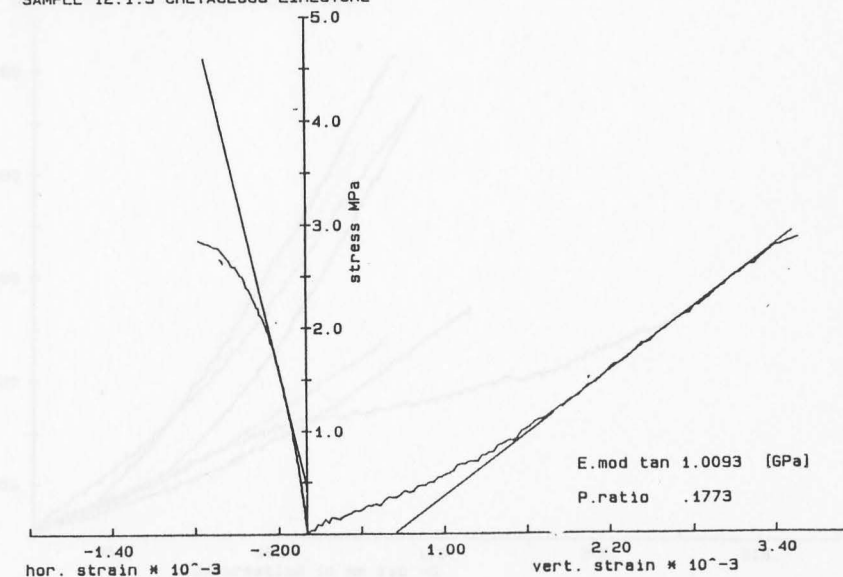
Laboratory Engineering Geology Delft University of Technology

SAMPLE 10.1.1 DEVONIAN LIMESTONE

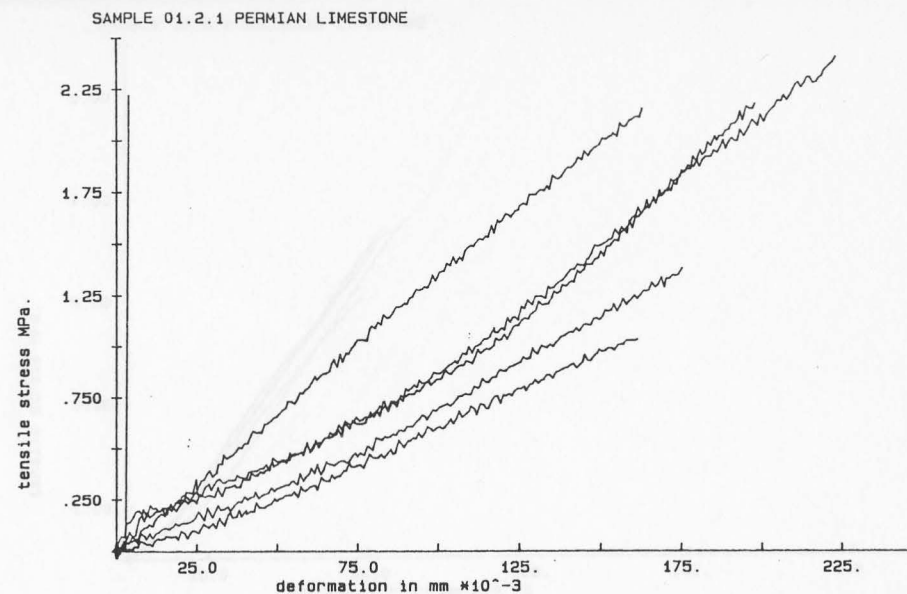
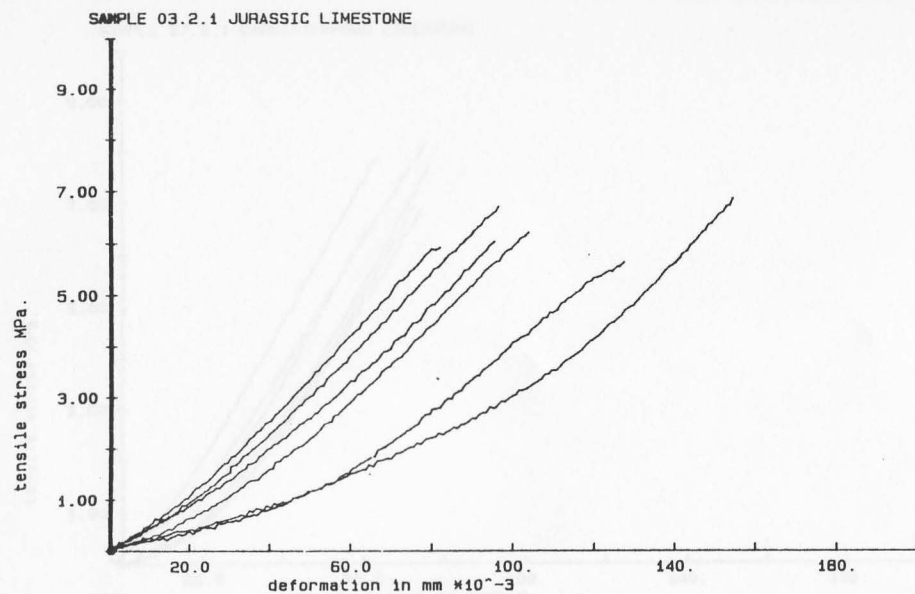


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SAMPLE 12.1.5 CRETACEOUS LIMESTONE

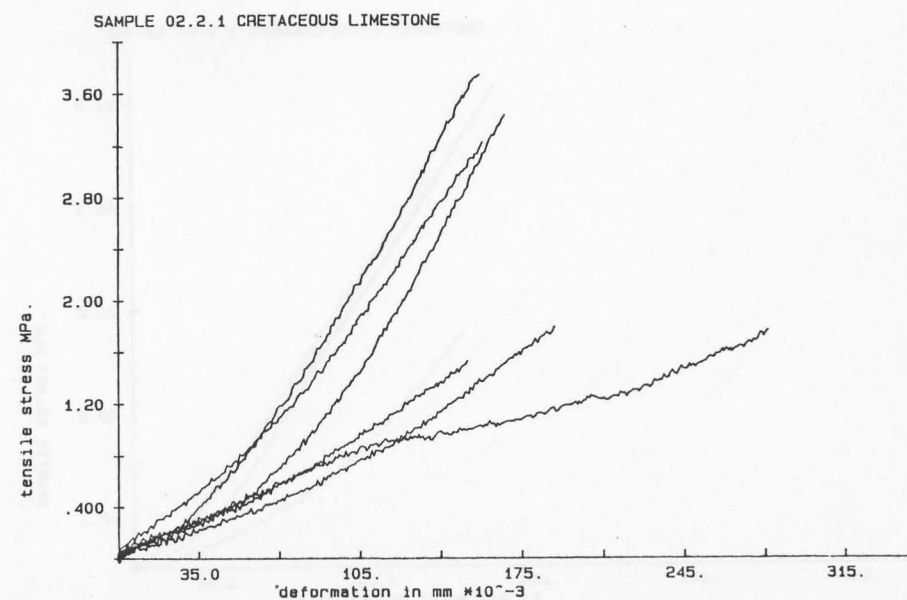
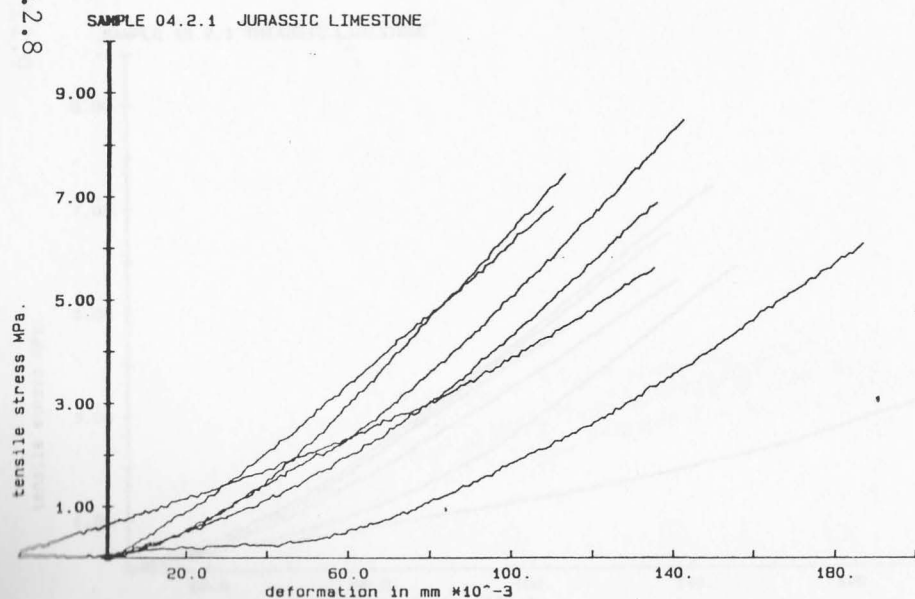


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Reference to literature cited within the text of this report



Reference to literature cited within the text of this report

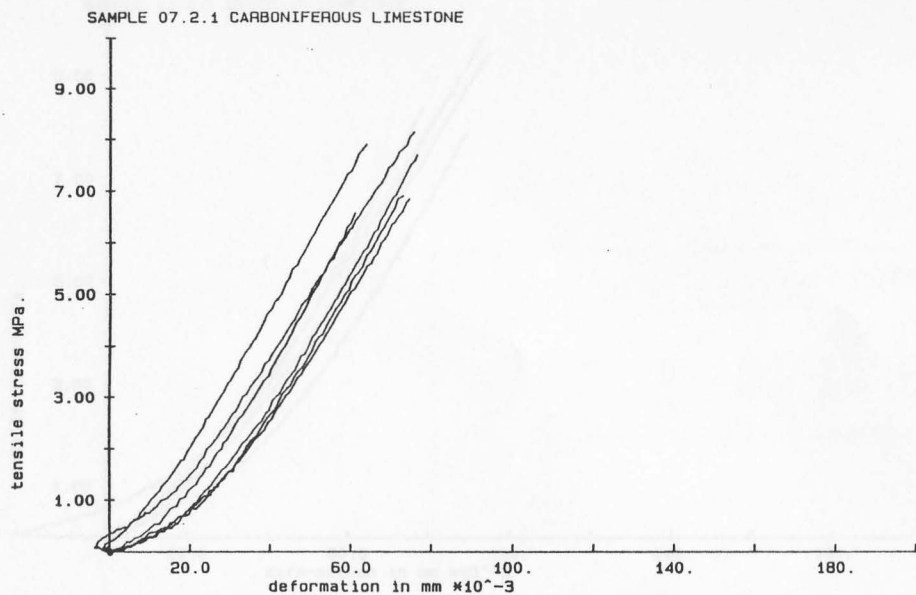


Reference to literature cited within the text of this report

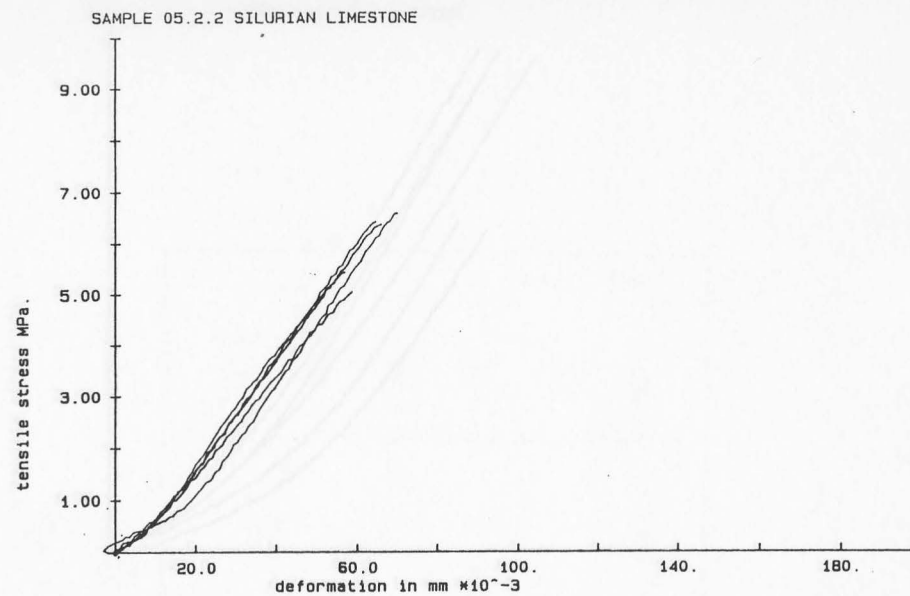


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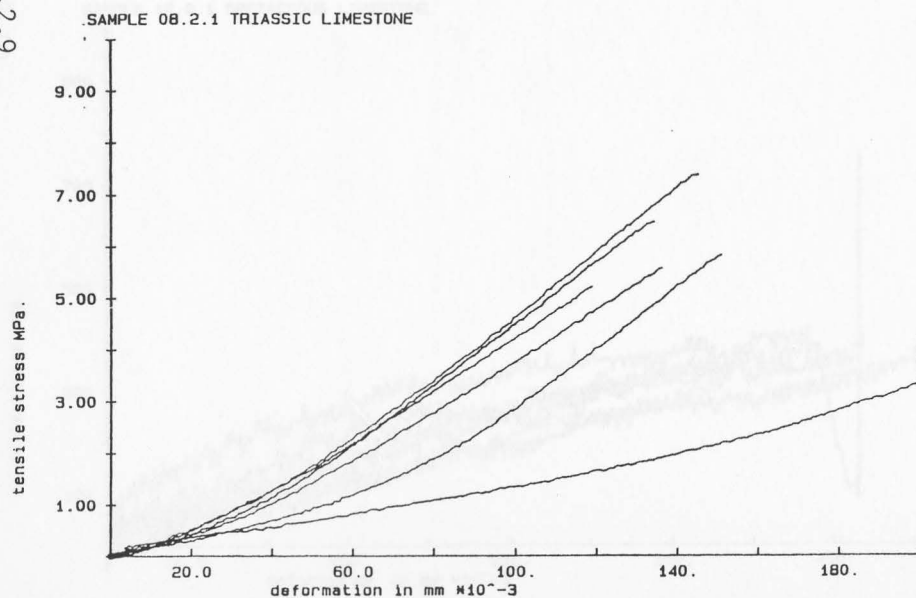




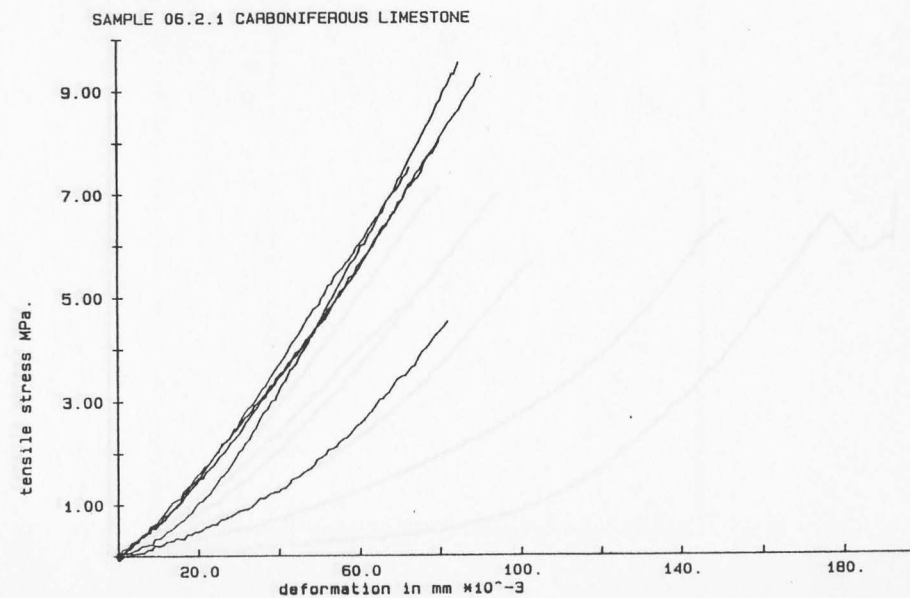
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Figure 1. Comparison of the calculated and experimental results for the case of the initial concentration of the solution is 0.1 g/g.



Figure 2. Comparison of the calculated and experimental results for the case of the initial concentration of the solution is 0.2 g/g.



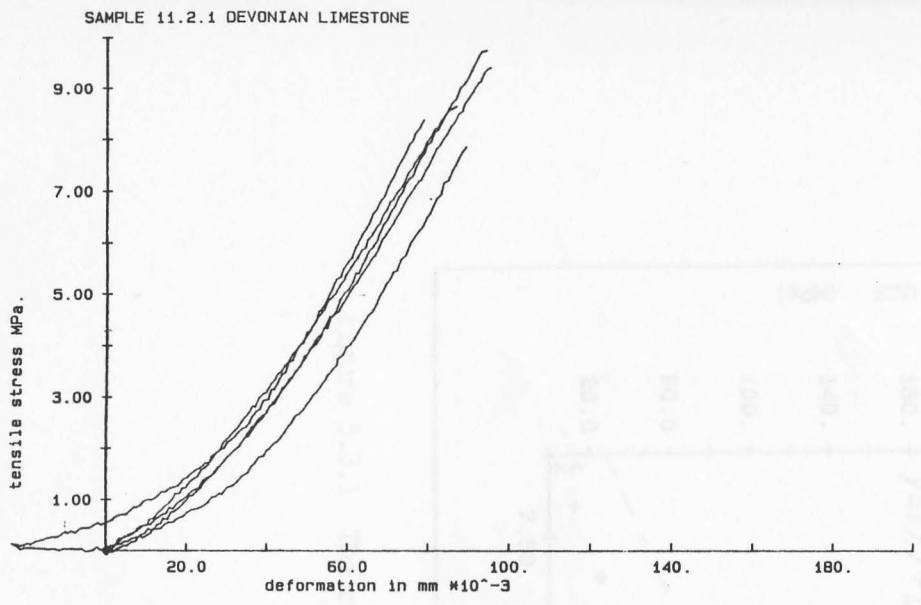
Figure 3. Comparison of the calculated and experimental results for the case of the initial concentration of the solution is 0.3 g/g.



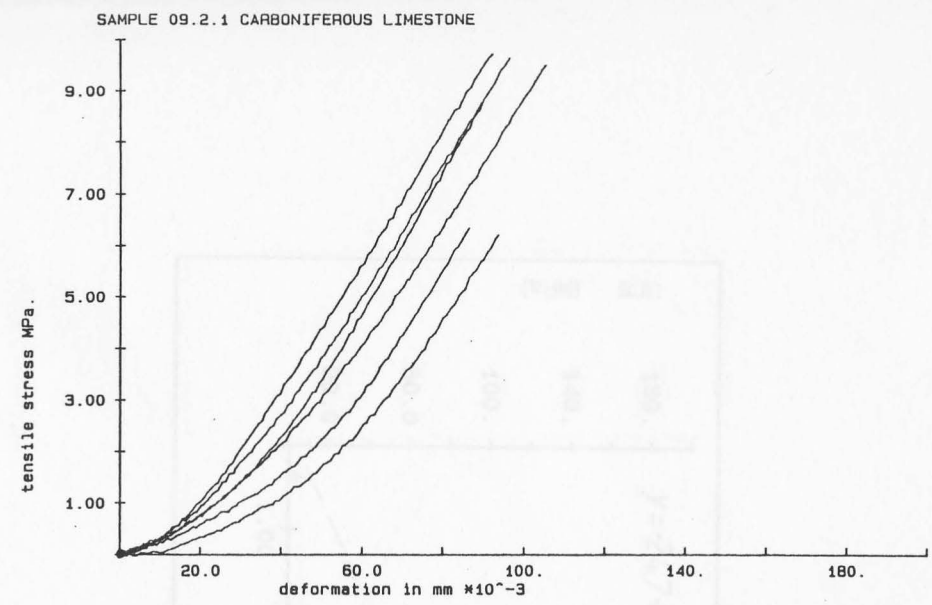
Figure 4. Comparison of the calculated and experimental results for the case of the initial concentration of the solution is 0.4 g/g.



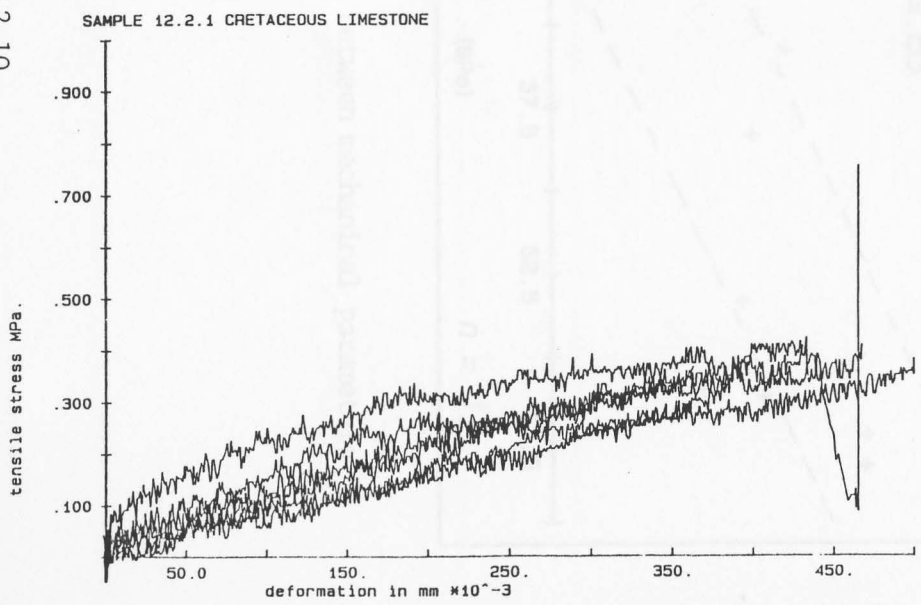
figure 5.2.10



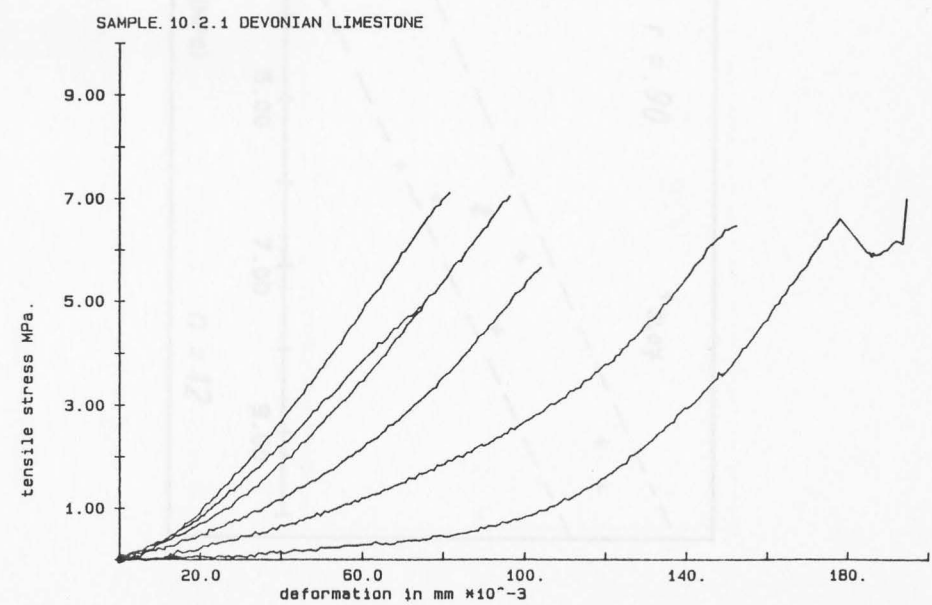
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CONCENTRATION OF POLYMER IN SOLUTION



Figure 1. Dependence of viscosity on concentration.

CONCENTRATION OF POLYMER IN SOLUTION



Figure 2. Dependence of viscosity on concentration.

CONCENTRATION OF POLYMER IN SOLUTION



Figure 3. Dependence of viscosity on concentration.

CONCENTRATION OF POLYMER IN SOLUTION



Figure 4. Dependence of viscosity on concentration.

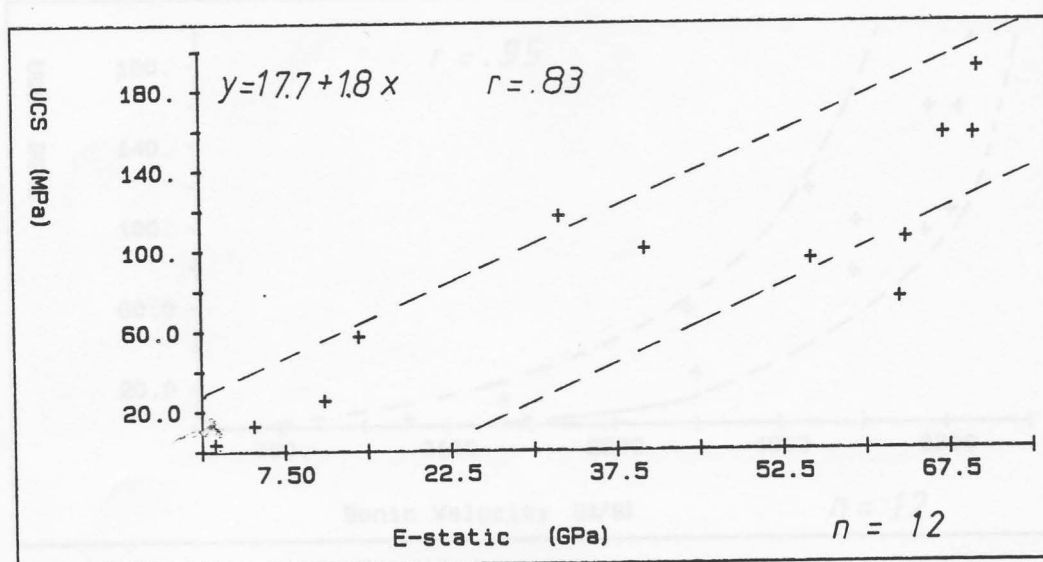
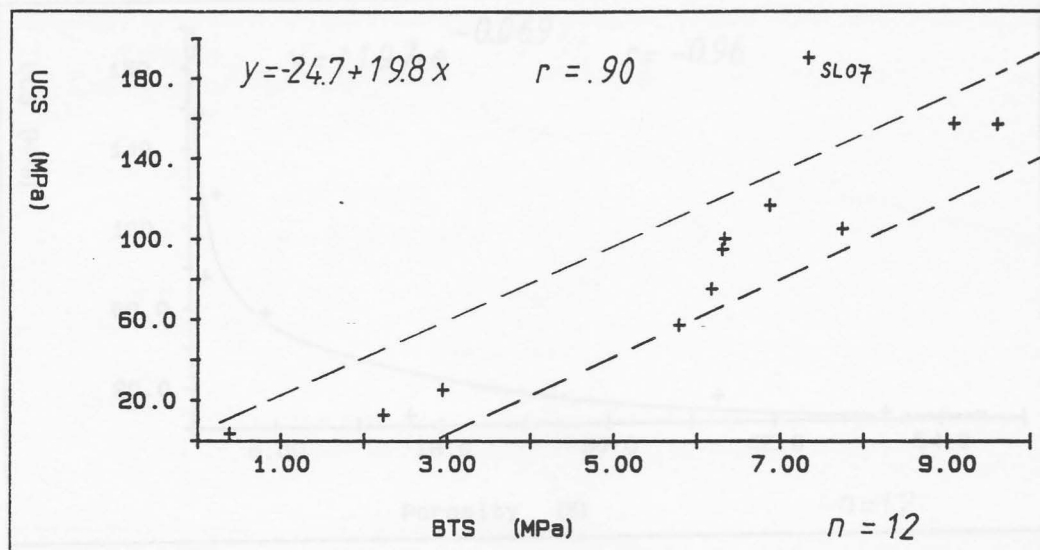


figure 5.3.1 The correlation between mechanical parameters

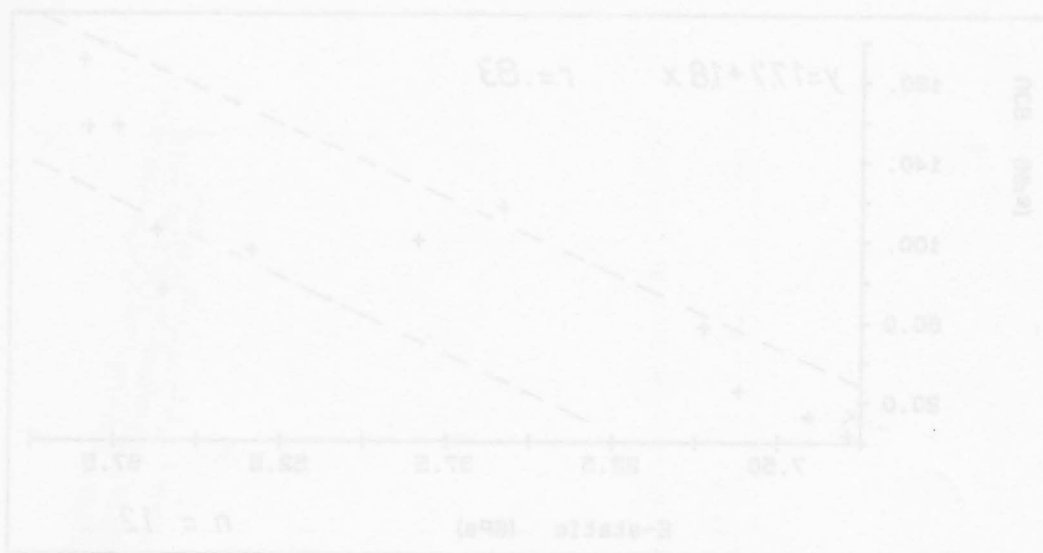
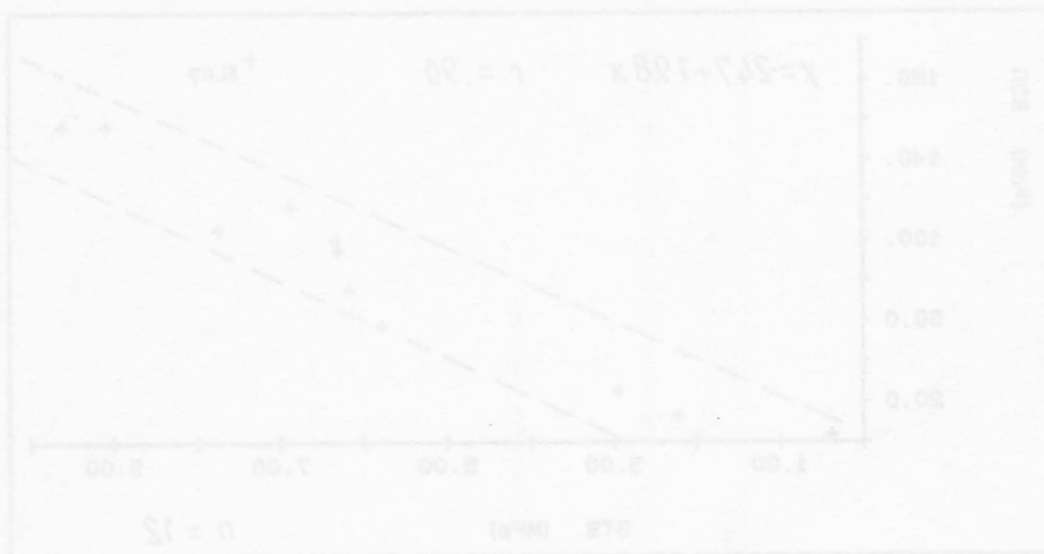


Figure 2.3.1 The correlation between mechanical parameters

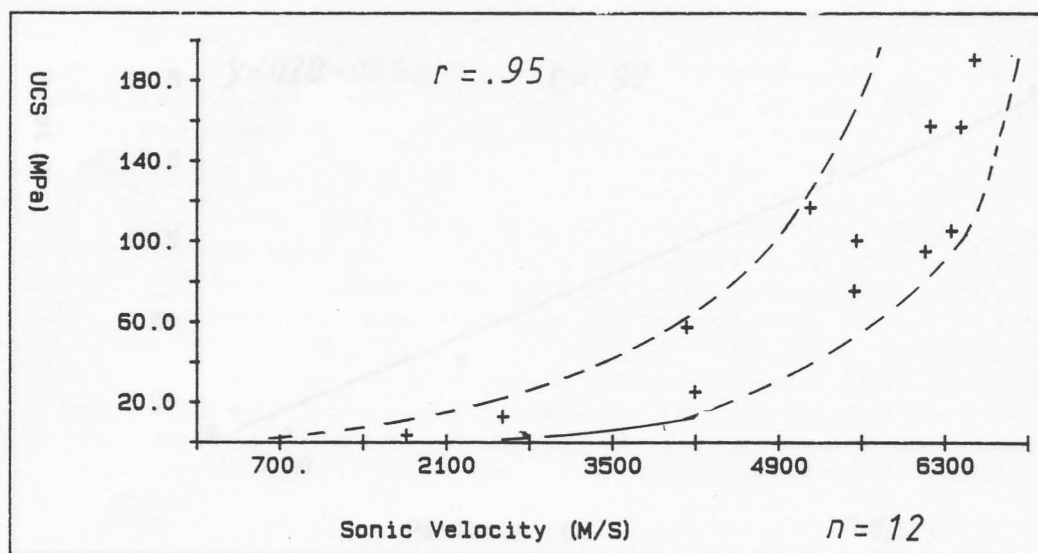
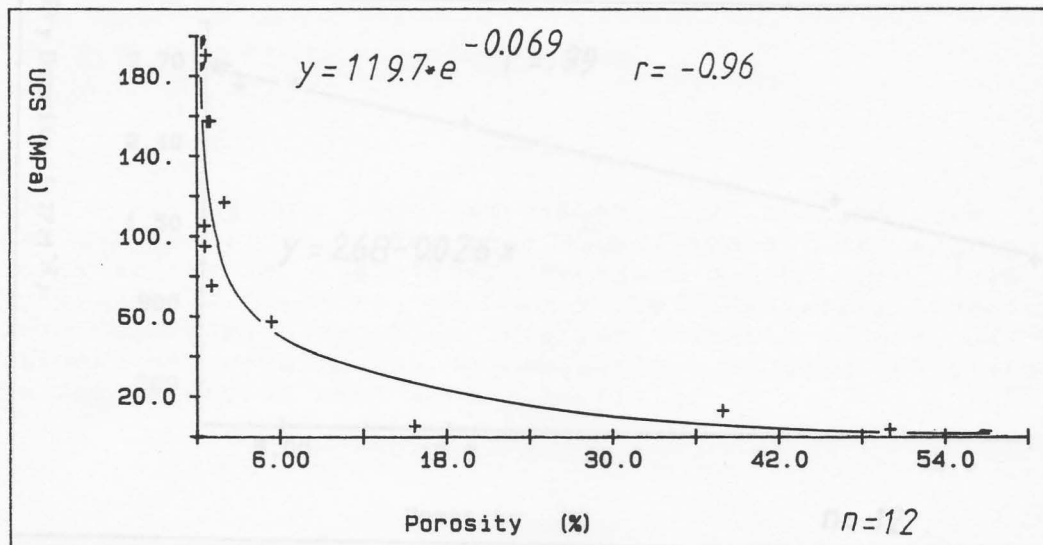


figure 5.3.2 The correlation between physical and mechanical parameters

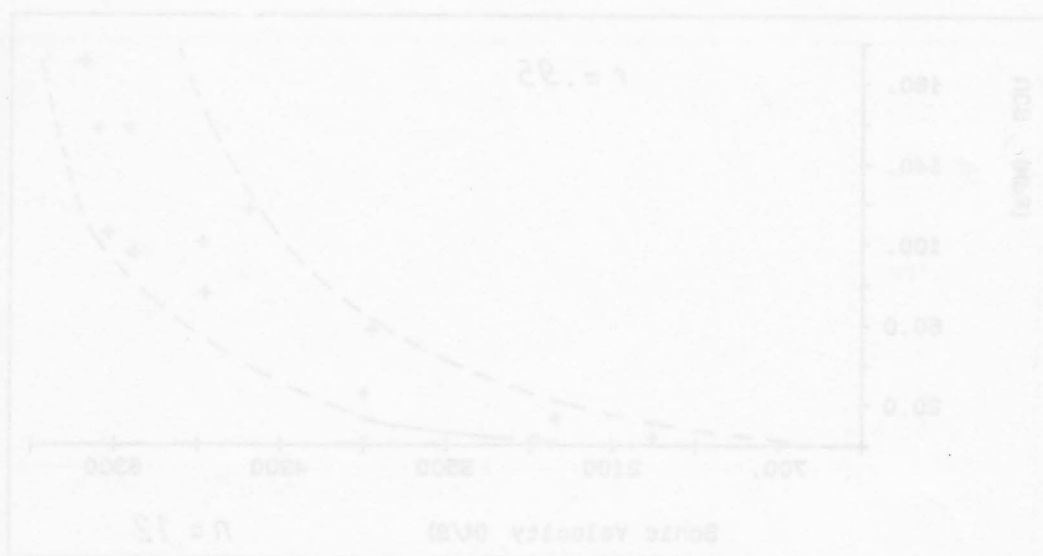
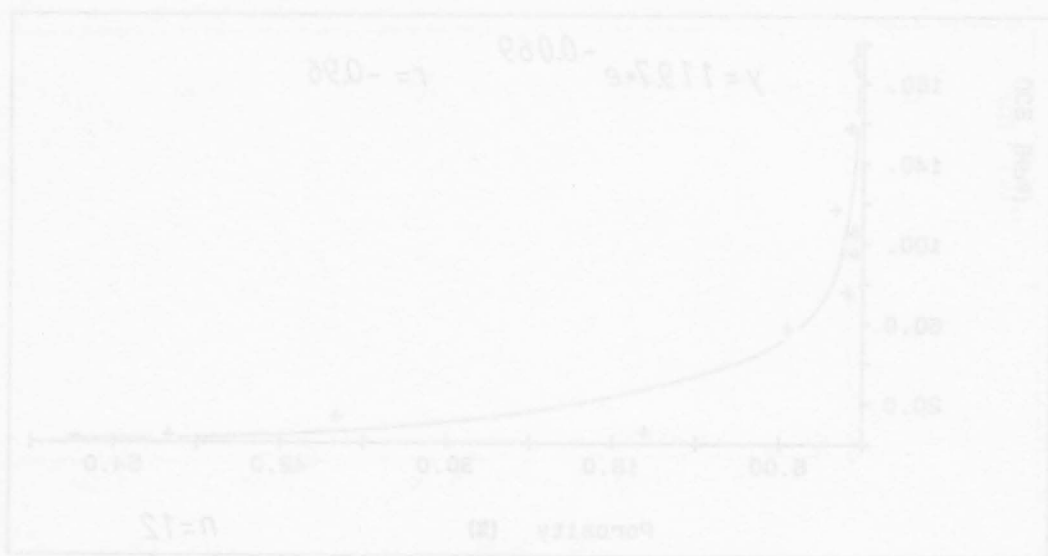


Figure 2.2.2. The correlation between physical and mechanical parameters

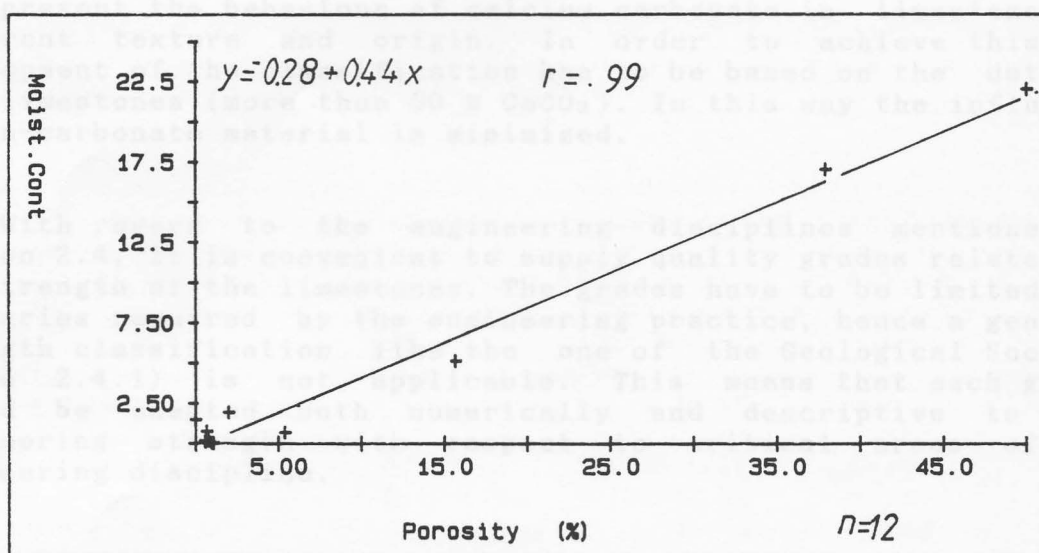
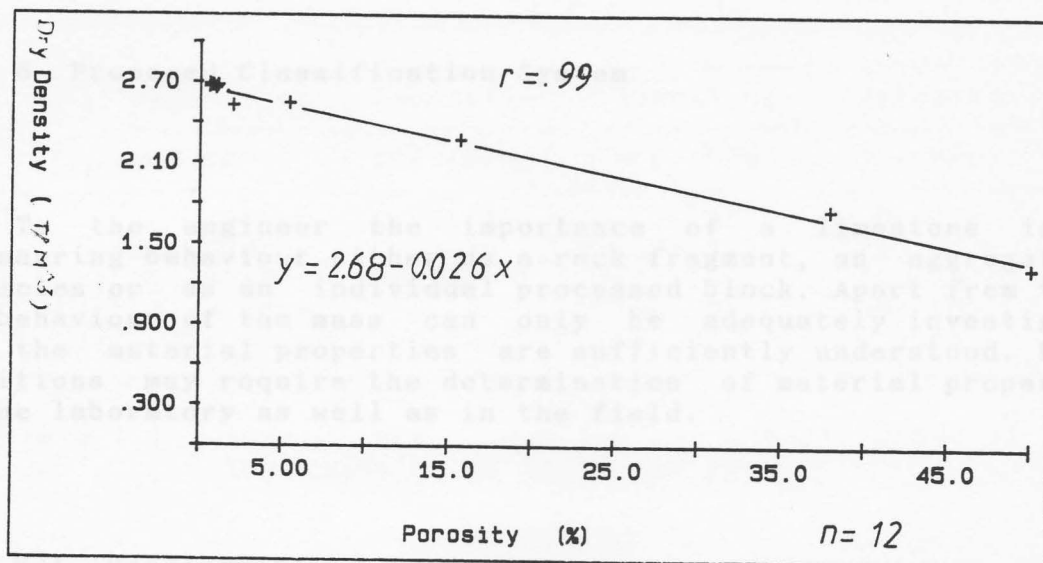


figure 5.3.4 The correlation between physical parameters

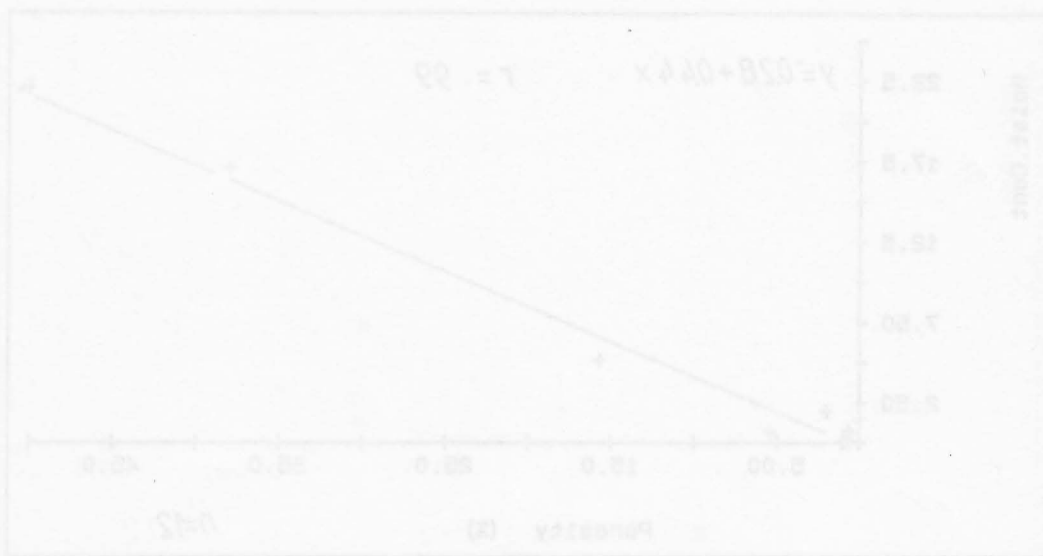
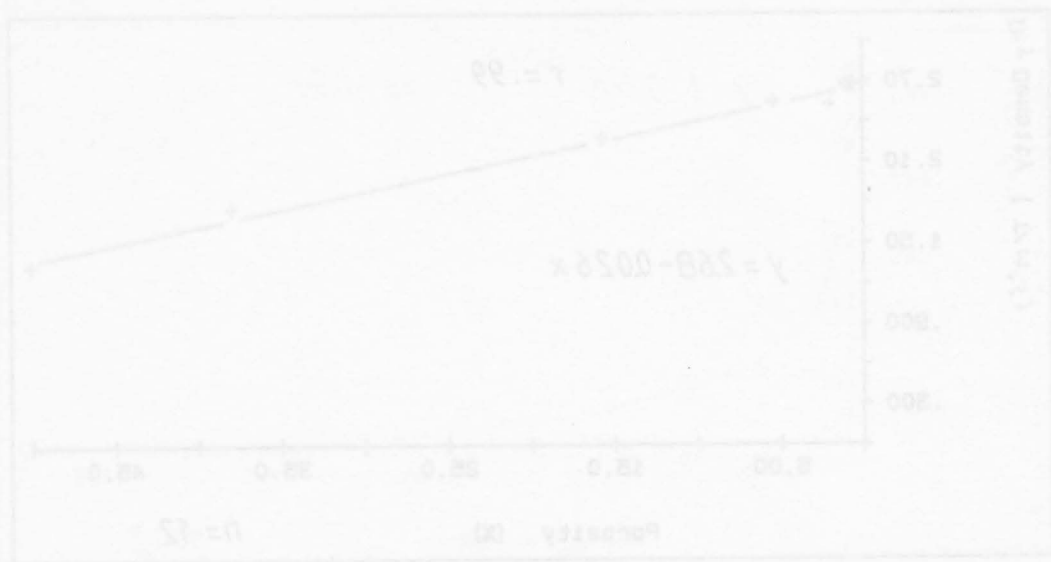


Figure 2.3.4. The correlation between physical parameters

6 Proposed Classification System

To the engineer the importance of a limestone is its engineering behaviour either as a rock fragment, an aggregate of particles or as an individual processed block. Apart from this, the behaviour of the mass can only be adequately investigated when the material properties are sufficiently understood. Hence conditions may require the determination of material properties in the laboratory as well as in the field.

6.1 Development

As concluded in section 2.4 a useful classification system for limestones should include the properties strength, density and grainsize of the intact rock. The relation between such properties may then form a fundamental basis for an empirical classification system. Nevertheless the essence of such a system must present the behaviour of calcium carbonate in limestones of different texture and origin. In order to achieve this the development of the classification has to be based on the data of pure limestones (more than 90 % CaCO_3). In this way the influence of non-carbonate material is minimized.

With regard to the engineering disciplines mentioned in section 2.4, it is convenient to supply quality grades related to the strength of the limestones. The grades have to be limited by boundaries required by the engineering practice, hence a general strength classification like the one of the Geological Society (table 2.4.1) is not applicable. This means that each grade should be adapted both numerically and descriptive to the engineering strength with respect to critical areas of the engineering discipline.

The empirical distinction of boundaries for the behaviour of rock material has been derived from the literature and if necessary modified to limestone circumstances. Below the four engineering disciplines are supplied with quality grades for Unconfined Compressive Strength (table 6.1.1).

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The empirical distinction of boundaries for the behavior of rock material has been derived from the literature and is necessary modified to limestone circumstances. Below the four engineering disciplines are supplied with quality grades for Unconfined Compressive Strength (Table 6.1.1).

Quality Grade	Excavation	Foundation	Tunneling	Aggregates
	UCS (MPa)	UCS (MPa)	UCS (MPa)	UCS (MPa)
I	> 100	0- 3	1- 25	0- 12
II	50 -100	3-10	25- 50	12- 25
III	12.5- 50	10-40	50-100	25- 38
IV	2.5- 12.5	40-80	100-200	38- 55
V	0- 2.5	> 80	> 200	> 55

table 6.1.1 Approximate ranges for the quality grade of limestones based on intact rock.

Source of ranges : Excavation Franklin (1971)
 Foundation CP for foundations
 (1972), Q30 (1984)
 Tunneling Bieniawski (1979)
 Aggregates ASTM C 33 (1979)

Additionally from the literature the following information was derived for limestone properties and procedures as a guidance to engineering practice (table 6.1.2).

Quality Grade	Excavation	Foundation	Tunnelling	Aggregates
	Method of removal	Safe Bearing Capacity (MPa)	Rate of Excavation m ³ /h per hp	Nature of optimal concrete
I	blast(fragment)	0-0.2	> 0.16	---
II	blast(fracture)	0.2-0.4	0.12-0.16	light
III	blast(loosen)	0.4-0.8	0.06-0.12	normal
IV	rip	0.8-2.0	0.03-0.06	normal
V	shovel	> 2.0	< 0.03	heavy

table 6.1.2 presumed values and descriptions of limestone qualities for engineering disciplines, under the assumption that unweathered and unfractured rock is involved.

To develop the classification system as described, the

Grade	Excavation	Foundation	Tunneling	Aggregate
UCS (MPa)	UCS (MPa)	UCS (MPa)	UCS (MPa)	UCS (MPa)
I	> 100	0-3	1-35	0-12
II	50-100	3-10	35-50	12-25
III	10-50	10-40	50-100	25-35
IV	2.5-12.5	40-80	100-200	35-55
V	0-2.5	> 80	> 200	> 55

Table 8.1.1. Approximate ranges for the quality grade of limestone based on intact rock.

Source of ranges :
 Excavation : Frensis (1971)
 Foundation : CP for Foundations (1972), QSD (1984)
 Tunneling : Rindfleisch (1978)
 Aggregate : ASTM C 33 (1978)

Additionally, from the literature the following information was derived for limestone properties and procedures as a guidance to engineering practice (Table 8.1.2).

Grade	Excavation	Foundation	Tunneling	Aggregate
UCS (MPa)	UCS (MPa)	UCS (MPa)	UCS (MPa)	UCS (MPa)
I	Plant (fragment)	0-0.2	> 0.15	light
II	Plant (fragment)	0.2-0.4	0.15-0.18	light
III	Plant (loose)	0.4-0.8	0.08-0.12	normal
IV	rip	0.8-2.0	0.03-0.08	normal
V	shovel	> 2.0	> 0.03	heavy

Table 8.1.2. Presumed values and descriptions of limestone qualities for engineering design. Under the assumption that unweathered and unfractured rock is involved.

To develop the classification system as described, the

exact relation of UCS versus Dry Density for pure limestone is a starting point (fig. 6.1.1). At certain angles to this e-power curve lines of equal CR-Index form the boundaries between the quality grades. These lines can be derived from the variation in the correlations between Dry Density and CR-Index, and UCS and CR-Index (fig. 6.1.2). The intervals at a constant CR-Index for UCS and Dry Density render three (x,y) pairs on the UCS-Dry Density plot. Through these three points the best fit represents the boundary (fig. 6.1.3). To incorporate the quality degrees in the diagram the intersecting points of the boundaries and the UCS-Dry Density curve are taken at the restricting strength's for each engineering discipline (fig. 6.1.4).

6.2 Mode of Operation (figure 6.2.1)

After selection of the required engineering discipline a descriptive classification renders a code based on the properties texture and grainsize. Additionally information about deformation modulus, abrasivity and cuttability could be supplied. Knowing the UCS and Dry Density of the limestone the quality grade can be determined. Finally, together with the descriptive classification code, the quality grade presents a measure of the engineering behaviour.

The application of this classification in the field can be realized with the help of light weight equipment; the Schmidt hammer for the UCS values; Hydrochloric acid (1.0 N) for the CaCO_3 content and the CR-Index; a lens for texture and grain size; a balance and a 100 ml cylinder and a small oven to measure the Dry Density. The volume of the limestone lump is then measured by the difference in the water height, when the lump is submerged.



exact relation of UCS versus Dry Density for pure limestone is a starting point (Fig. 8.1.1). At certain angles to this upper curve lines of equal CB-Index form the boundaries between the quality grades. These lines can be derived from the variation in the correlations between Dry Density and CB-Index, and UCS and CB-Index (Fig. 8.1.2). The intervals at a constant CB-Index for UCS and Dry Density render three (x,y) pairs on the UCS-Dry Density plot. Through these three points the best fit represents the boundary (Fig. 8.1.3). To incorporate the quality degrees in the diagram the intersecting points of the boundaries and the UCS-Dry Density curve are taken as the starting strength for each engineering discipline (Fig. 8.1.4).

8.2 Mode of Operation (Figure 8.2.1)

After selection of the required engineering discipline a descriptive classification renders a code based on the properties texture and grain size. Additionally information about deformation modulus, abrasivity and collectibility could be supplied. Knowing the UCS and Dry Density of the limestone the quality grade can be determined. Finally, together with the descriptive classification code, the quality grade presents a measure of the engineering behaviour.

The application of this classification in the field can be realized with the help of light weight equipment; the Schmidt hammer for the UCS values; Hydrochloric acid (1.5 N) for the CaCO₃ content and the CB-Index; a lens for texture and grain size; a balance and a 100 ml cylinder and a small oven to measure the Dry Density. The volume of the limestone lump is then measured by the difference in the water height, when the lump is submerged.

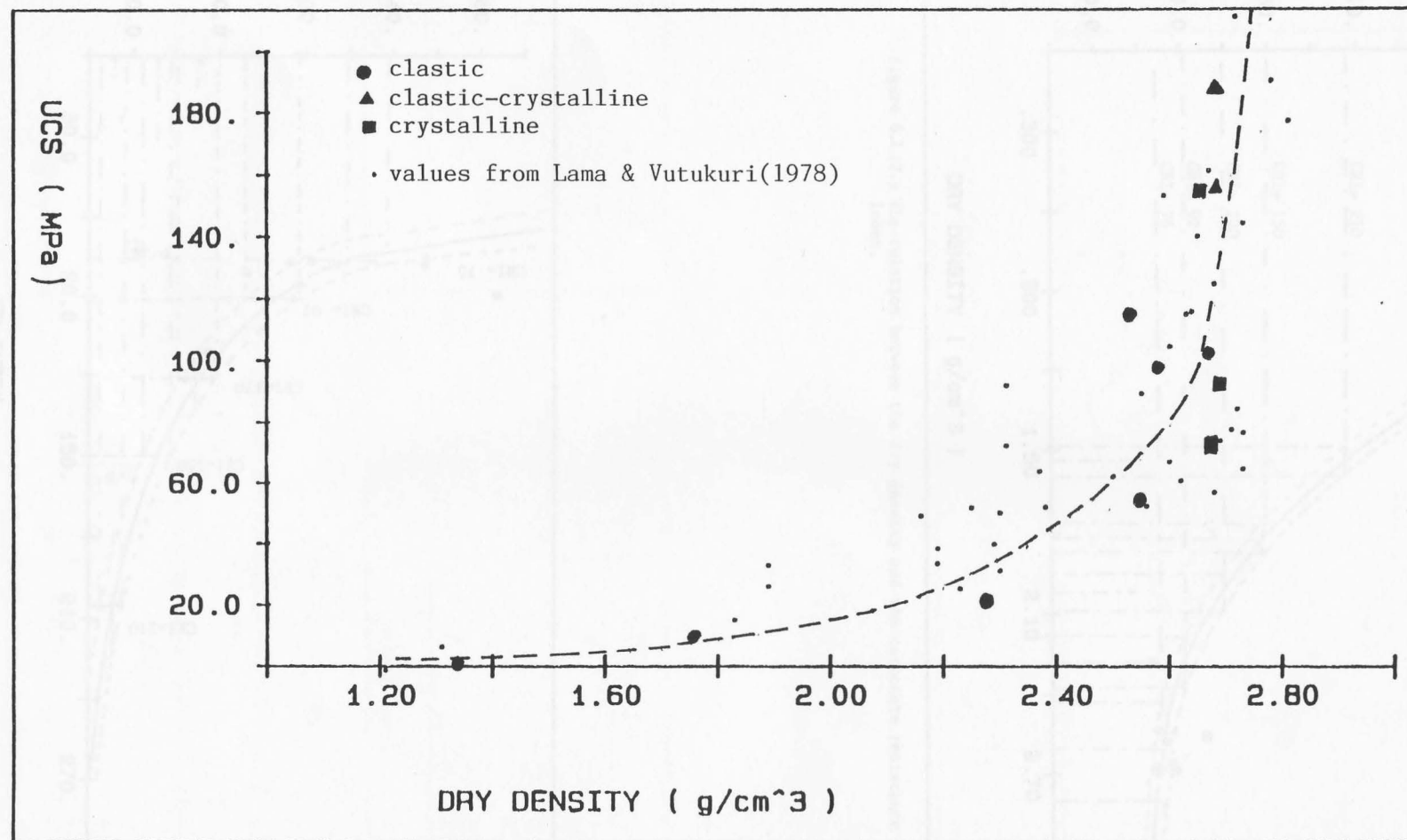


figure 6.1.1 The relation between dry density and strength for limestones.

Figure 1. Relationship between the initial and final values of the parameter α for the polymer.



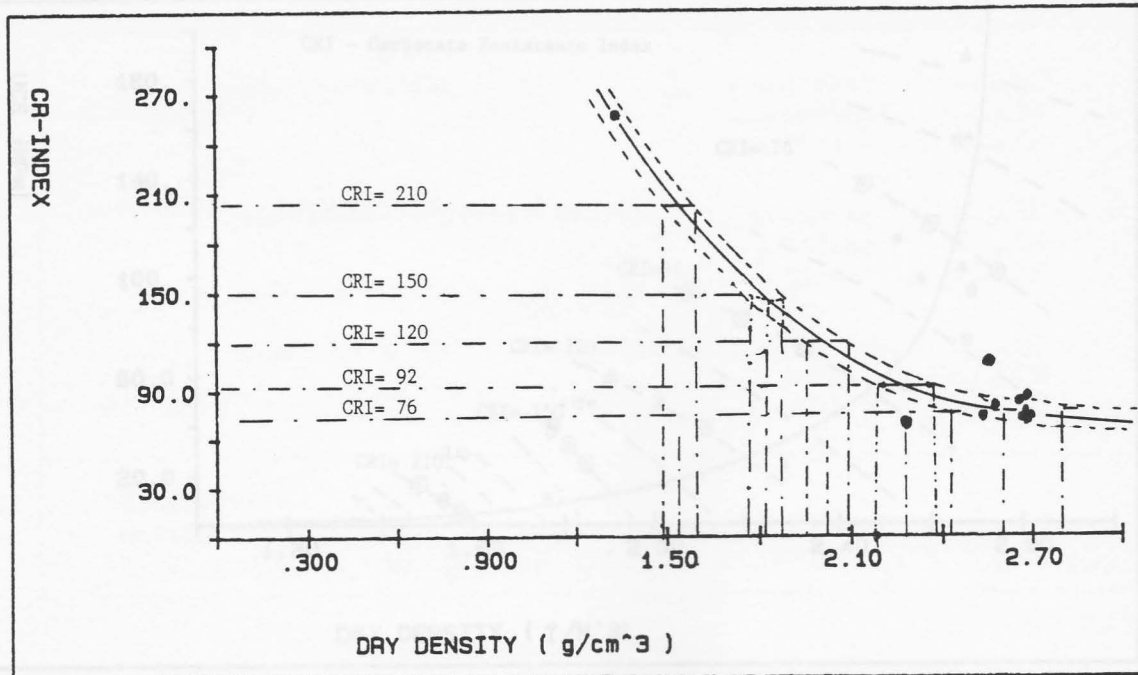


figure 6.1.2.a The relation between the dry density and the carbonate resistance Index.

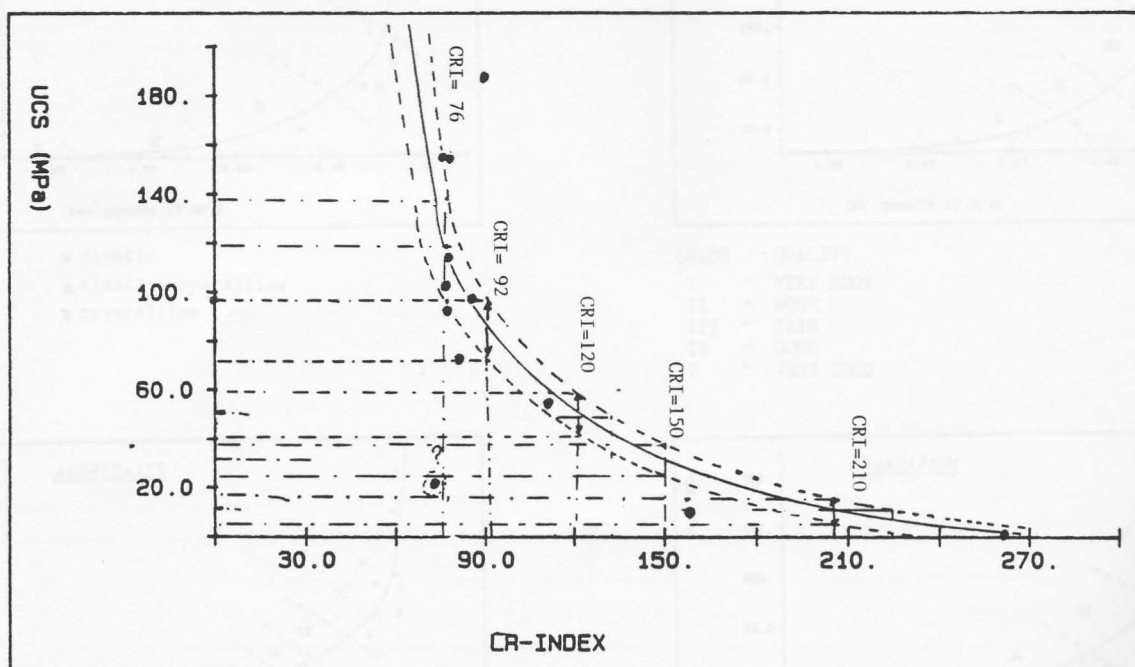


figure 6.1.2.b The relation between the unconfined compressive strength and the carbonate resistance index.

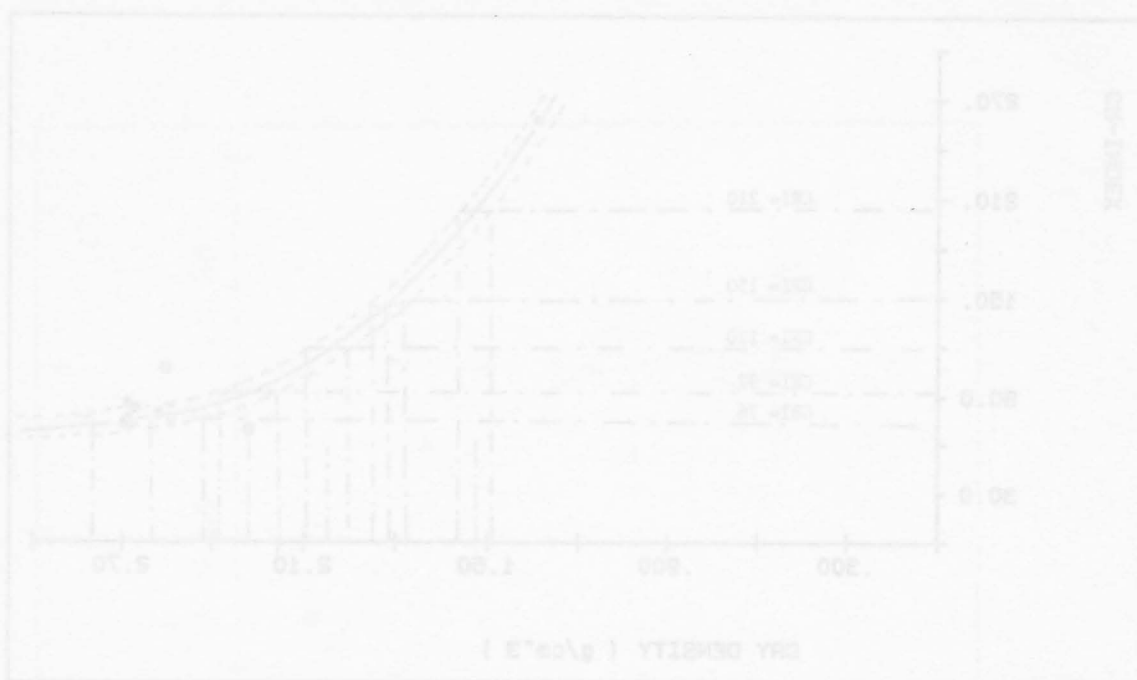


Figure 5.1.1: The relation between the dry density and the CR-Index.

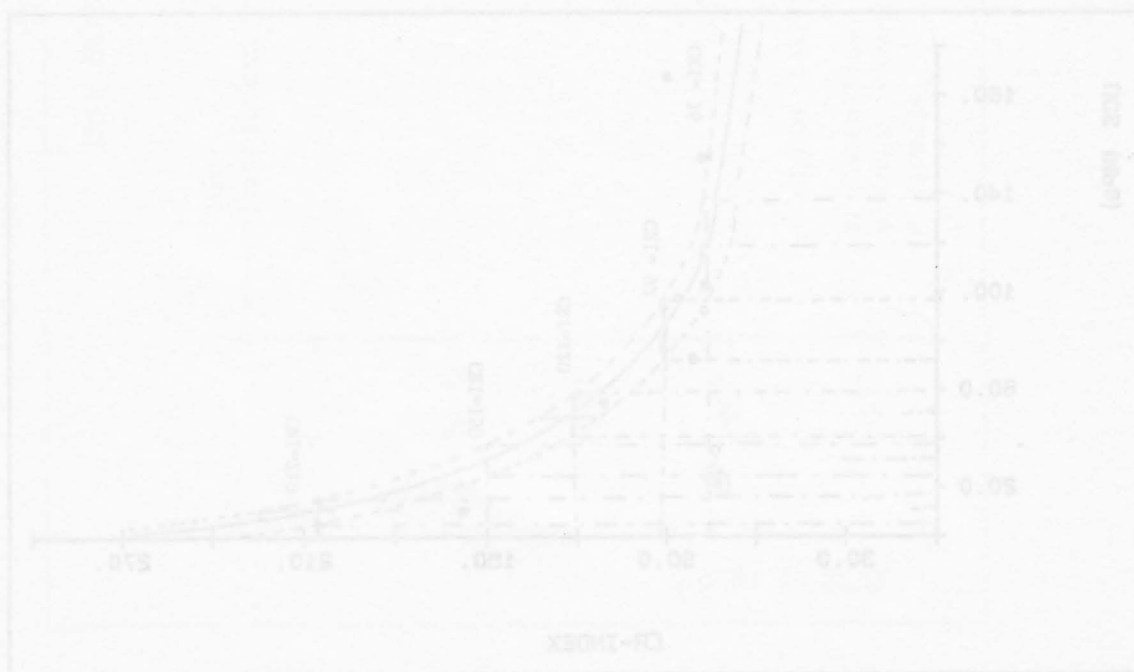


Figure 5.1.2: The relation between the CR-Index and the CR-Value.

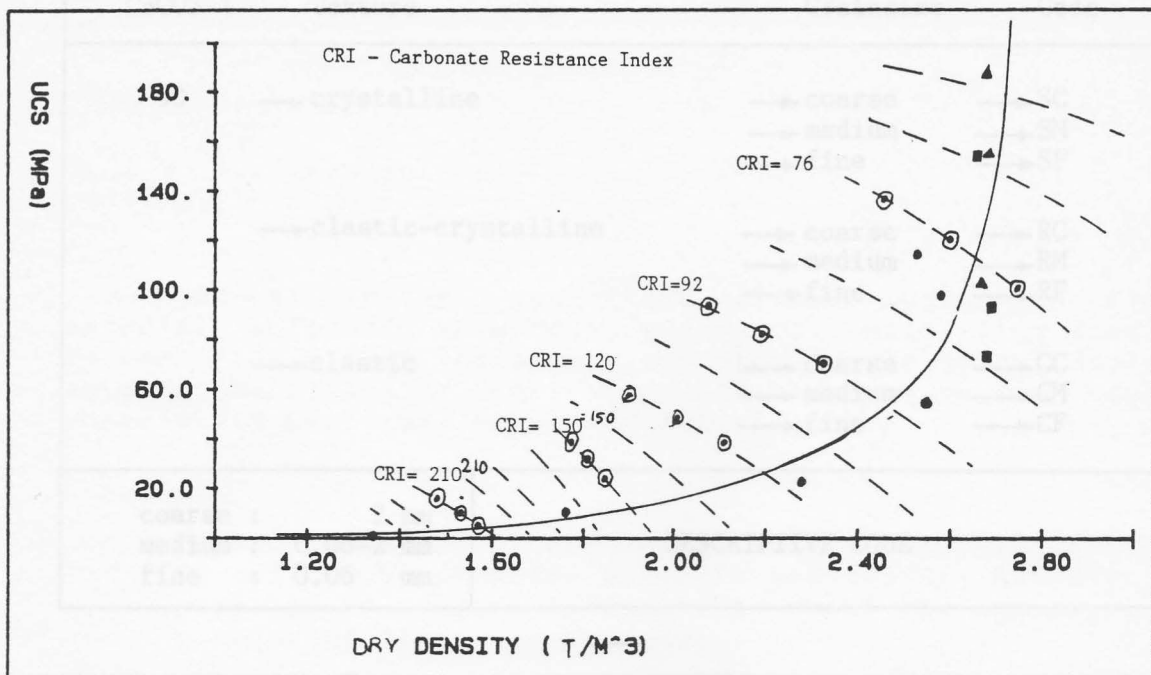
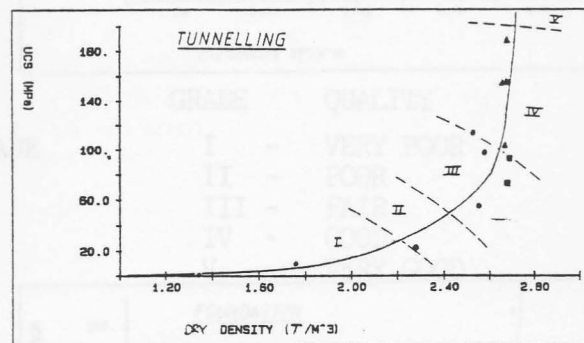
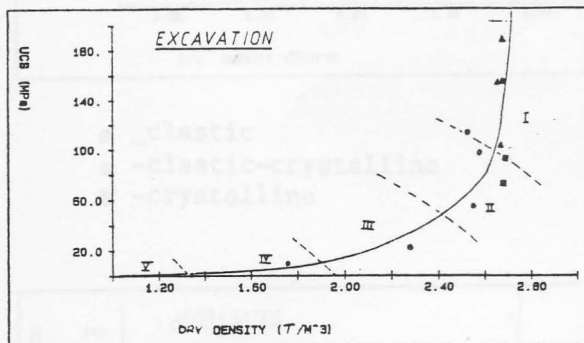


figure 6.1.3 Boundaries of constant CRI incorporated for the UCS-Dry Density relation.



GRADE QUALITY

I - VERY POOR

II - POOR

III - FAIR

IV - GOOD

V - VERY GOOD

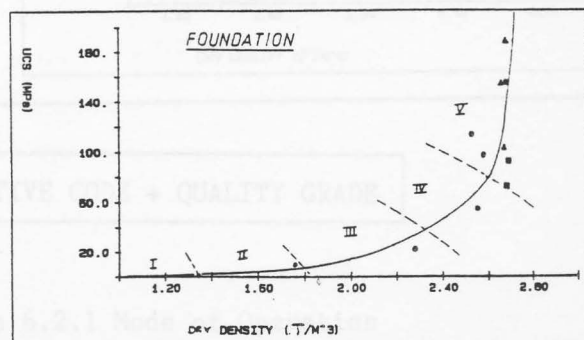
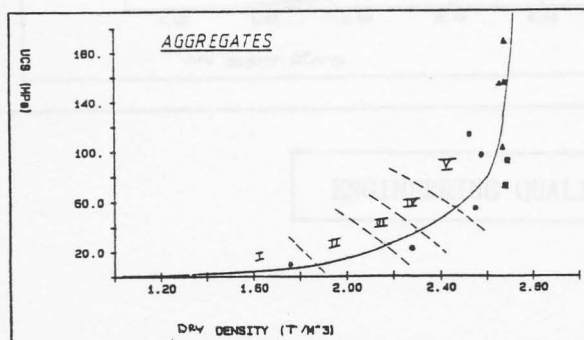


figure 6.1.4 Quality degrees for limestone material in engineering disciplines.

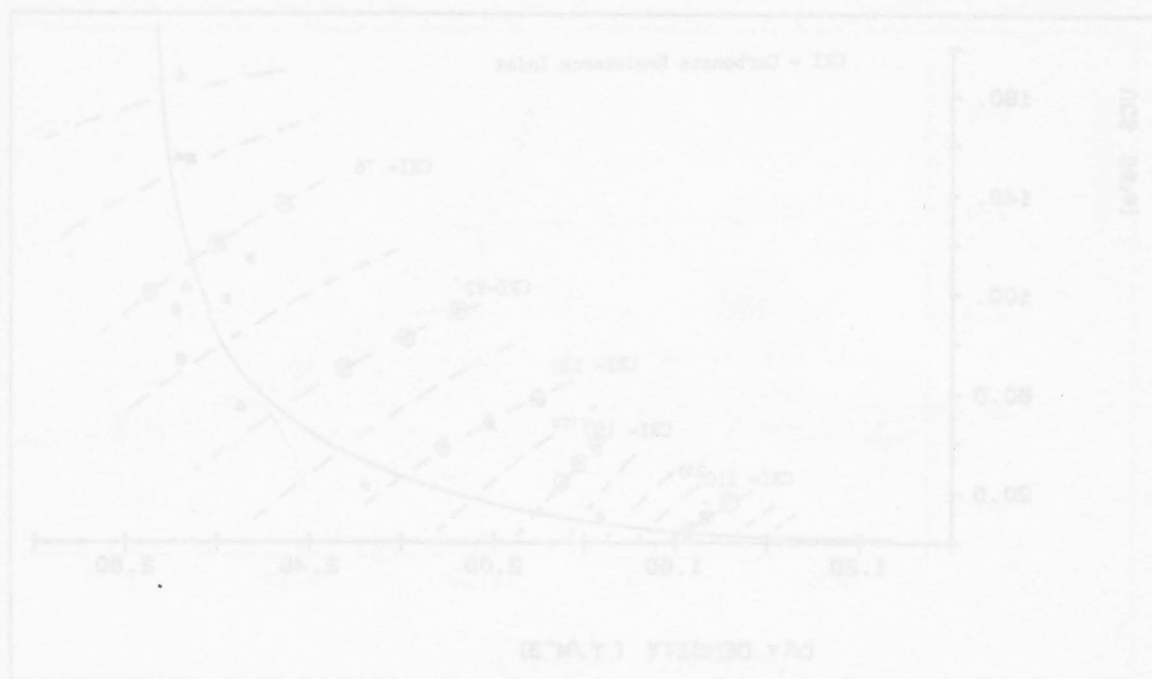
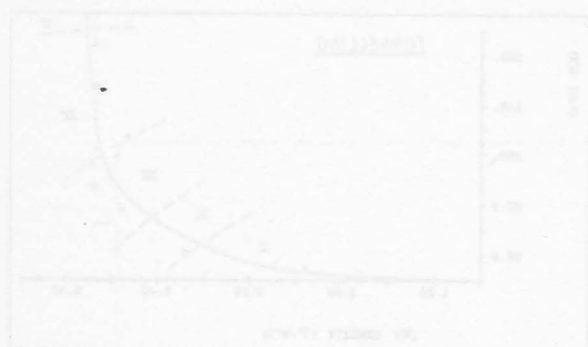
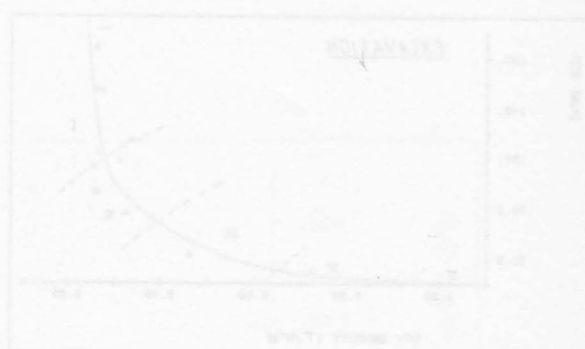


Figure 5.15: Plot of Dry Density (T.M.B) vs. Unit Weight (T.M.B) for various CBI specimens.



Specimen	Dry Density (T.M.B)	Unit Weight (T.M.B)
CBI-10	1.50	50.0
CBI-15	1.60	60.0
CBI-20	1.70	70.0
CBI-25	1.80	80.0
CBI-30	1.90	90.0
CBI-35	2.00	100.0



a classic
a classic
a classic

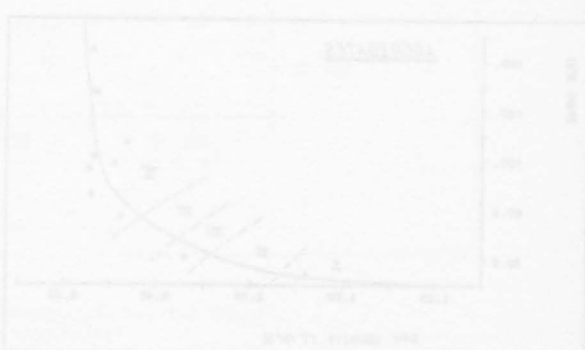
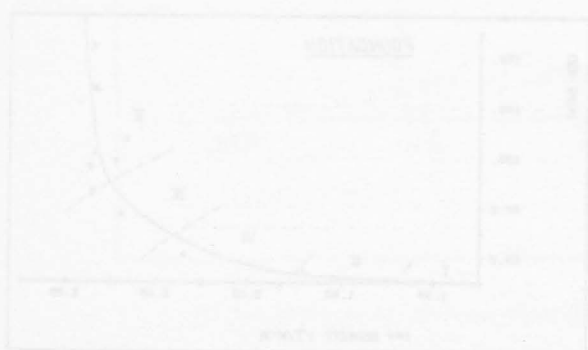
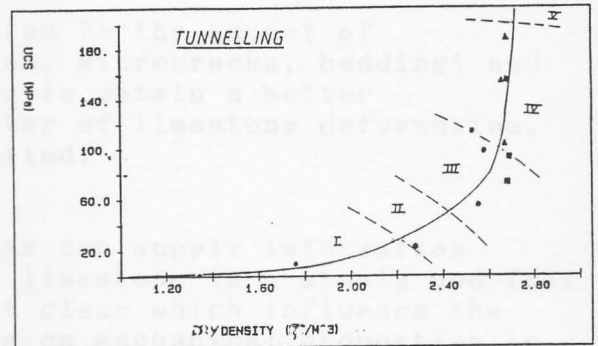
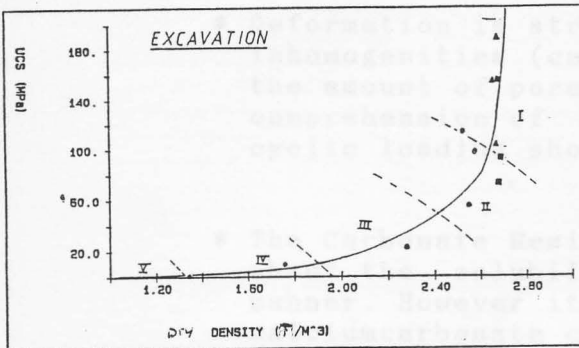


Figure 5.16: Plot of Dry Density (T.M.B) vs. Unit Weight (T.M.B) for various CBI specimens.

Classification of Limestone material
for engineering purposes

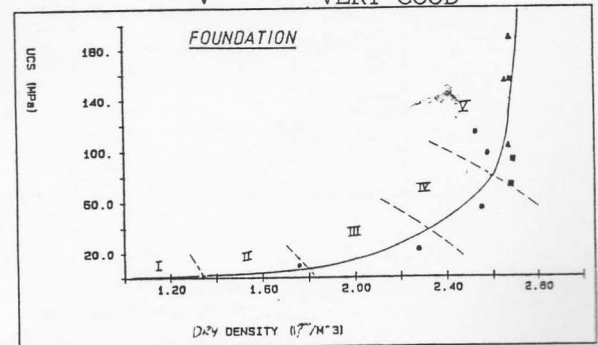
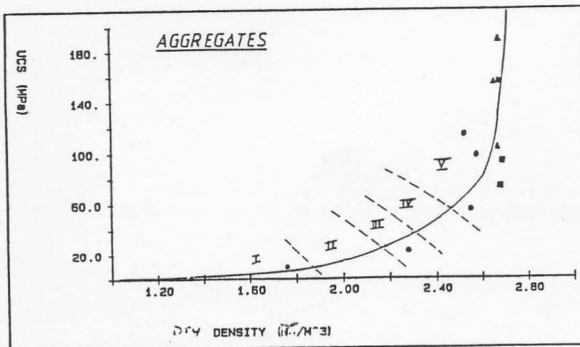
CaCO3 %	Texture	Grainsize	Code
90	→ crystalline	→ coarse	→ SC
		→ medium	→ SM
		→ fine	→ SF
	→ clastic-crystalline	→ coarse	→ RC
		→ medium	→ RM
		→ fine	→ RF
	→ clastic	→ coarse	→ CC
		→ medium	→ CM
		→ fine	→ CF
coarse : 2 mm		DESCRIPTIVE CODE	
medium : 0.06-2 mm			
fine : 0.06 mm			



- -clastic
- ▲ -clastic-crystalline
- -crystalline

QUALITY GRADE

GRADE	QUALITY
I	- VERY POOR
II	- POOR
III	- FAIR
IV	- GOOD
V	- VERY GOOD

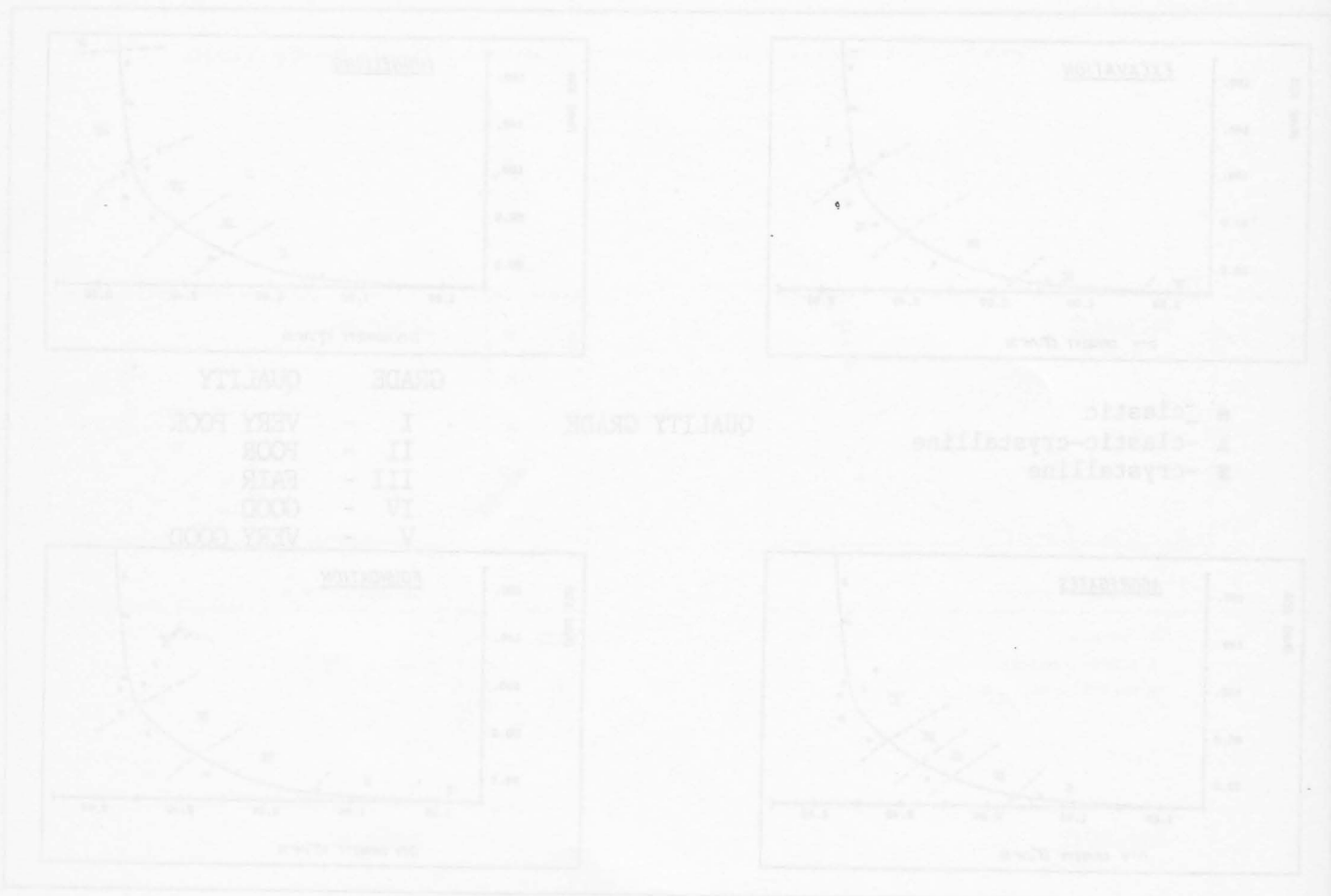


ENGINEERING QUALITY= DESCRIPTIVE CODE + QUALITY GRADE

figure 6.2.1 Mode of Operation

Classification of limestone material
for engineering purposes

Grain size	Texture	Grade
coarse medium fine	crystalline	10
coarse medium fine	crystalline-crystalline	11
coarse medium fine	crystalline	12
coarse medium fine	crystalline	13
coarse medium fine	crystalline	14
coarse medium fine	crystalline	15
coarse medium fine	crystalline	16
coarse medium fine	crystalline	17
coarse medium fine	crystalline	18
coarse medium fine	crystalline	19
coarse medium fine	crystalline	20
coarse medium fine	crystalline	21
coarse medium fine	crystalline	22
coarse medium fine	crystalline	23
coarse medium fine	crystalline	24
coarse medium fine	crystalline	25
coarse medium fine	crystalline	26
coarse medium fine	crystalline	27
coarse medium fine	crystalline	28
coarse medium fine	crystalline	29
coarse medium fine	crystalline	30
coarse medium fine	crystalline	31
coarse medium fine	crystalline	32
coarse medium fine	crystalline	33
coarse medium fine	crystalline	34
coarse medium fine	crystalline	35
coarse medium fine	crystalline	36
coarse medium fine	crystalline	37
coarse medium fine	crystalline	38
coarse medium fine	crystalline	39
coarse medium fine	crystalline	40
coarse medium fine	crystalline	41
coarse medium fine	crystalline	42
coarse medium fine	crystalline	43
coarse medium fine	crystalline	44
coarse medium fine	crystalline	45
coarse medium fine	crystalline	46
coarse medium fine	crystalline	47
coarse medium fine	crystalline	48
coarse medium fine	crystalline	49
coarse medium fine	crystalline	50
coarse medium fine	crystalline	51
coarse medium fine	crystalline	52
coarse medium fine	crystalline	53
coarse medium fine	crystalline	54
coarse medium fine	crystalline	55
coarse medium fine	crystalline	56
coarse medium fine	crystalline	57
coarse medium fine	crystalline	58
coarse medium fine	crystalline	59
coarse medium fine	crystalline	60
coarse medium fine	crystalline	61
coarse medium fine	crystalline	62
coarse medium fine	crystalline	63
coarse medium fine	crystalline	64
coarse medium fine	crystalline	65
coarse medium fine	crystalline	66
coarse medium fine	crystalline	67
coarse medium fine	crystalline	68
coarse medium fine	crystalline	69
coarse medium fine	crystalline	70
coarse medium fine	crystalline	71
coarse medium fine	crystalline	72
coarse medium fine	crystalline	73
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coarse medium fine	crystalline	81
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coarse medium fine	crystalline	84
coarse medium fine	crystalline	85
coarse medium fine	crystalline	86
coarse medium fine	crystalline	87
coarse medium fine	crystalline	88
coarse medium fine	crystalline	89
coarse medium fine	crystalline	90
coarse medium fine	crystalline	91
coarse medium fine	crystalline	92
coarse medium fine	crystalline	93
coarse medium fine	crystalline	94
coarse medium fine	crystalline	95
coarse medium fine	crystalline	96
coarse medium fine	crystalline	97
coarse medium fine	crystalline	98
coarse medium fine	crystalline	99
coarse medium fine	crystalline	100



ENGINEERING QUALITY - DESCRIPTIVE CODE + QUALITY GRADE

Figure 3.2.1 Mode of Operation

7 Recommendations

The engineering geological classification of limestone material is founded upon requirements of engineering disciplines on one side and on geological features on the other. Having this in mind the following conclusions can be based on the results of experiments performed on intact material :

- * From the empirical determined relation between the Unconfined Compressive Strength and the Dry Density it is possible to derive information about the engineering quality of limestone material.
- * Deformation is strongly related to the amount of inhomogenities (calcite veins, microcracks, bedding) and the amount of pores. In order to obtain a better comprehension of the character of limestone deformation, cyclic loading should be applied.
- * The Carbonate Resistance Index can supply information about the solubility of the limestone in a simple and fast manner. However it is not yet clear which influence the calciumcarbonate content has on mechanical properties in relation to other constituents. Hence impure carbonates cannot be classified in the same manner.

Fig. 4.1 Suggested sample description sheet

Appendix 4.1 Suggested sample description sheet for carbonate rocks
(Burnett & Spas 1979)

7 Recommendations

The engineering geological classification of limestone material is founded upon requirements of engineering classification on one side and on geological features on the other. Having this in mind the following conclusions can be based on the results of experiments performed on tested material:

1. From the empirical determined relation between the Uniaxial Compressive Strength and the Dry Density it is possible to derive information about the engineering quality of limestone material.

2. Deformation is strongly related to the amount of inhomogeneities (cavities, veins, microcracks, bedding) and the amount of pores. In order to obtain a better comprehension of the character of limestone deformation, cyclic loading should be applied.

3. The Carbonate Resistance Index can supply information about the variability of the limestone in a single and last manner. However, it is not yet clear which influence the calcareous content has on mechanical properties in relation to other constituents. Hence, further research cannot be classified in the same manner.

Fig. 6. A suggested standardised sample description sheet

Appendix A.2 The description of rocks for engineering purposes -material (IAEG 1981)

3. Description of Rocks für Engineering Purposes

Description of rock involves the following steps:

- (i) Determination of the fundamental rock name: the 'lithological rock name';
- (ii) Description of the properties of the rock material;
- (iii) Description of additional properties necessary to describe the features of the rock mass.

The properties of the rock mass are controlled partly by the properties of the rock material, but in many rock masses structural features substantially control the engineering properties. Such features include structures and discontinuities such as joints and bedding plane partings, and the distribution of rock and soil materials in the weathering profile.

The three steps listed above provide a 'descriptive rock name' from which engineering properties may more readily be inferred than from a 'lithological rock name'.

3.1 The Descriptive Rock Name

In a rock description the main characteristics should be given in the following order:

Rock name
 Supplementary petrographic properties
 Rock material properties
 Colour
 Texture
 Grain size
 Other textural features and fabric
 State of weathering
 State of alteration
 Strength
 Rock Mass Properties
 Structure
 Discontinuities
 Weathering profile

The descriptive scheme has been modified from that recommended in Anon. (1972b). The main differences are in the treatment of state of weathering and the weathering profile, and an expansion in the description of structure. Structural aspects have been dealt with more thoroughly by Anon. (1977) and also in I.S.R.M. (1977).

3.1.1 The Lithological Name: The lithological rock name is of primary importance because it indicates the genetic rock group and provides basic information on mineral composition and grain size. Supplementary petrographic properties may be used where necessary to qualify the rock name, signifying for example a relative abundance of a particular mineral — biotite granite — or indicating minor admixtures of other lithological types. These supplementary features may be extremely useful as a means of discriminating between different rocks that have the same lithological name. Minor constituents may also have an important effect on the mechanical and physical properties of rocks, and should be carefully considered.

The rock name is selected from the classification tables (Table 1) and these are the only rock names that are recommended for use. In arriving at a name for a rock, there is no substitute either for geological knowledge or for an aid to identification that is reliable and easy to apply.

LIGHTNESS	CHROMA	HUE
Light	pinkish	pink
Dark	reddish	red
	yellowish	yellow
	brownish	brown
	greenish	green
	bluish	blue
		white
	greyish	grey
		black

Table 2: Terms for lightness, chroma and hue which may be used in combination for colour description

3.2 Description of Rock Material

3.2.1 Colour: Rock colour can be quantitatively evaluated using, for example, the Rock Color Chart published by the Geological Society of America (Anon. 1963). As an alternative it is recommended that the following simple system (Anon. 1972b), which serves to limit the subjectivity of an estimation, should be used. One term is selected, as required, from each column (Table 2), and combined as a colour assessment.

Examples of use are: light yellowish brown, dark reddish brown, dark brown, etc. If necessary colour differences can be emphasized separately by the use of terms such as spotted, dappled, mottled, streaked, for example light yellowish brown spotted with dark brown.

3.2.2 Texture: Of the textural elements used for description and classification, the most important is grain size which, for the predominant size of grain, can be classified semiquantitatively. From Table 1 the relations between rock names and grain sizes can be understood. It will be recalled that the class boundaries have been fixed at limits of grain size grades adopted for engineering soils, that is the boundaries between clay, silt and sand sizes that are justified and determined by the differences in the physical behaviour of those soils (Glossop and Skempton, 1945).

Because grain size considerably affects the physical properties of a rock it should always be indicated directly in the rock description rather than relying on the grain size implication in the rock name.

It is usually sufficient to estimate grain size by eye, which may be aided by a hand lens in the case of fine-grained and amorphous rocks. The limit of unaided vision is approximately 0.06 mm.

Many other aspects of rock texture may be used to amplify the description, such as:

3.2.2.1 Relative grain size: for example uniform, non-uniform, porphyritic

3.2.2.2 Grain shape: may be described by reference to the general form of the particles, their angularity which indicates the degree of rounding at edges and corners, and their surface characteristics (Table 3).

FORM	equidimensional
	flat
	elongated
	flat and elongated
	irregular
ANGULARITY	angular
	subangular
	subrounded
	rounded
SURFACE CHARACTERISTICS	rough
	smooth

Table 3: Terms used in the description of grain shape

3.2.2.3 Fabric: the spatial arrangement of grains in the rock may show a preferred orientation or lack of it, and may produce patterns by non-uniform arrangements of grains, crystals and groundmass.

3.2.2.4 Porosity: the size, shape, orientation of pore or void spaces should be described.

3.2.3 State of Weathering: Description of the state of weathering of rock material is of particular importance in describing engineering rocks because weathering has profound effects on the physical and mechanical properties of rock material. In any description there needs to be a statement whether or not the rock material is considered to be either in a fresh state or is weathered. Weathering effects may be described in terms of discoloration, chemical decomposition or physical disintegration.

The extent of particular weathering effects may be sub-divided using such qualifying terms, for example 'highly decomposed', 'extremely discoloured', 'slightly disintegrated', as will aid the description of the material being examined. These descriptive qualifying terms may be quantified if necessary by estimation from drill core or in the natural exposure (Table 4).

TERM	DEGREE OF CHANGE (per cent)
Fresh	0
Slightly	Over 0 - 10
Moderately	10 - 35
Highly	35 - 75
Extremely	Over 75

Table 4: Terms for the description of the degree of weathering of rock material

Depending on the character and distribution of the weathering changes, and the extent to which a rigid rock framework is retained, the weathered rock material may assume the characteristics of an engineering soil at an early stage.

Extremely weathered rock material will almost certainly be an engineering soil, and may be classed as a residual soil if the original rock fabric has collapsed or changed so as to remove most traces of the original fabric.

Examples of use are: fresh rock; slightly decomposed; moderately disintegrated; highly discoloured. Usually combinations occur: highly disintegrated and moderately decomposed, etc.

3.2.4 State of Alteration: The terms used for weathering of rock material may be used where appropriate as in many instances the effects of alteration may not be easily distinguishable from those brought about by weathering. Wherever possible common terms should be used, e. g. slightly kaolinised, highly mineralised; the terms may be quantified using the scale in Table 4.

3.2.5 Strength: The uniaxial compression test gives a reliable indication of the strength of rock material, although the test results are dependent on the moisture content of the specimen, any anisotropy in the material, and the test procedure adopted. A scale of strength is given in Table 5.

As an alternative method of strength testing for use in the field, the point load test (Broch and Franklin, 1972; ISRM, 1977) is recommended. Provided that individual test results are normalised to a standard specimen thickness of 50 mm, and the recommended test procedures

TERM	COMPRESSIVE STRENGTH (MPa)
Weak	1.5*- 15
Moderately strong	15 - 50 ⁺
Strong	50 - 120
Very strong	120 - 230
Extremely strong	Over 230

*Rocks with a strength under 1.25 MPa are, as a rule, hard soils and should be tested accordingly

⁺Soft rocks are weaker than 50 MPa; strong rocks are stronger than 50 MPa

Table 5: A scale of strength for dry rock material

are followed, this test provides a good estimate of unconfined compressive strength. The relation:

$$UCS = 25 PLS$$

where UCS is the unconfined compressive strength and PLS is the point load strength, has been demonstrated repeatedly and can be accepted as a reasonably reliable approximation.

The piston-press test, devised by Srejner, Petrova and Jakusev (1958) and described by Matula (1969), is a quick method of determining the strength and deformation properties of rock materials. Test values show a very close correlation with the results of the standard unconfined compressive strength performed on cubes of rock.

DETRITAL SEDIMENTARY					CHEMICAL/ ORGANIC	GENETIC GROUP		
BEDDED						Usual Structure		
Grains of rock, quartz, feldspar and clay minerals			At least 50% of grains are of carbonate		Salts, carbonates, silica, carbonaceous	Composition		
RUDACEOUS	Grains are of rock fragments		Limestone (undifferentiated)	Calci- rudite	SALINE ROCKS Halite Anhydrite Gypsum	Very coarse- grained	60	PREDOMINANT GRAIN SIZE (mm)
	Rounded grains: CONGLOMERATE Angular grains: BRECCIA					Coarse- grained		
AREN- ACEOUS	Grains are mainly mineral fragments		Limestone (undifferentiated)	Calc- arenite	CALCAREOUS ROCKS LIMESTONE DOLOMITE	Medium- grained	2	
	SANDSTONE Grains are mainly mineral fragments					Fine- grained		
ARGILLACEOUS or LUTACEOUS	MUDSTONE SHALES: fissile mudstone	SILTSTONE 50% fine- grained particles	Marlstone	Calci- siltite	SILICEOUS ROCKS Chert Flint CARBONACEOUS ROCKS LIGNITE COAL	0.06	0.002	
		Claystone 50% very fine grained particles		CHALK Calci- lutite		Very fine- grained		
					SILICEOUS ROCKS	GLASSY AMORPHOUS		

Table 1: Classification of rock types: sedimentary

Table 1: Classification of rock types: sedimentary

TERM	COMPRESSIVE STRENGTH (MPa)
Weak	1.37 - 15
Weakly strong	15 - 30
Strong	30 - 120
Very strong	120 - 250
Extremely strong	Over 250

Units with a strength index of 15 MPa and above are classified as strong and should be tested accordingly.

Units with a strength index of 15 MPa and above are classified as strong and should be tested accordingly.

Table 1. A scale of strength for the rock masses.

are known, this can provide a good estimate of rock mass strength. The equation:

$$UCS = 22 P_L$$

where UCS is the unconfined compressive strength and P_L is the point load strength, has been demonstrated to be a good estimate of rock mass strength.

The point load test, devised by Brown, Brown and Johnson (1970) and described by Brown (1970) is a quick method of determining the strength and deformation properties of rock materials. The test gives a very close correlation with the results of the uniaxial compressive strength test, but is much easier to perform in the field.

SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
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SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS		METAMORPHIC		IGNEOUS	
SEDIMENTARY		METAMORPHIC											

Appendix B.1 Porosity Determination with the "Ruska" Universal Porositymeter following the method of Kobe

Apparatus

The universal porositymeter consists of; a mercury pump; with on top the pyknometer barrel, which can be opened to insert a specimen; above the barrel a needle valve, which is used to close the barrel when pressure builds up; and two manometers, one for high pressure and one for low pressure.

Principle

The porosity measurement with the method of Kobe enables us to determine the porosity of a small kernel in a faster way. The method of Kobe is based on Boyle's law, which says that under a constant temperature, the product of pressure and volume of an ideal gas is also constant. If two reference volumes are taken at 40cc and at 30cc, which are then compressed to 30 Psi. From Boyle's law it follows that :

$$\begin{aligned} 40.P_a &= V_f.P_r \\ 30.P_a &= V_e.P_r \end{aligned} \quad \begin{aligned} & \frac{P_r}{P_r - P_a} = \frac{10}{10 - V_e - V_f} \end{aligned} \quad (1)$$

The volume of the material (V_g) is calculated from :

$$\begin{aligned} P_a.(V_f - V_g) &= P_r.(V_f' - V_g) \\ V_g &= \frac{P_r}{P_r - P_a} . (V_f' - V_f) \end{aligned} \quad (2)$$

were P_a = the atmospheric pressure
 P_r = Pressure at 30 Psi.
 V_r = reference volume of uncompressed air
 V_f = volume of 40cc air after compression
 V_f' = volume of 40cc air and kernel after compression
 V_e = volume of 30cc air after compression

when equation (1) is substituted in equation (2) one finds :

Appendix B.1 Potentially Determination with the "Rocks"
Universal Potentiometer following the method of
Kohn

Apparatus

The universal potentiometer consists of: a mercury pump;
with on top the pyrometer barrel, which can be opened to insert
a specimen; above the barrel a needle valve, which is used to
close the barrel when pressure builds up; and two manometers, one
for high pressure and one for low pressure.

Principle

The porosity measurement with the method of Kohn enables us
to determine the porosity of a small kernel in a faster way. The
method of Kohn is based on Boyle's law, which says that under a
constant temperature, the product of pressure and volume of an
ideal gas is also constant. If two reference volumes are taken at
40cc and at 30cc, which are then compressed to 30 Psi from
Boyle's law it follows that:

$$\begin{aligned} 40 P_1 &= V_1 P_2 \\ 30 P_1 &= V_2 P_2 \end{aligned} \quad (1)$$

The volume of the material (V_2) is calculated from:

$$\begin{aligned} P_2 (V_1 - V_2) &= P_1 (V_1 - V_2) \\ V_2 &= \frac{P_1 (V_1 - V_2)}{P_2 - P_1} \end{aligned} \quad (2)$$

where P_1 = the atmospheric pressure
 P_2 = pressure at 30 Psi
 V_1 = reference volume of uncompressed air
 V_2 = volume of 40cc air after compression
 V_1' = volume of 30cc air and kernel after compression
 V_2' = volume of 30cc air after compression

when equation (1) is substituted in equation (2) one finds:

$$V_g = \frac{10}{10 - V_e - V_f} \cdot (V_f' - V_f) \quad (3)$$

the porosity is then calculated from : $\Phi = 1 - \frac{V_g}{V_b}$ (4)

V_b = bulk volume

Appendix B.2 Permeability measurement

Liquid permeameter

the liquid permeameter is an apparatus which uses pressurized water to create a flow through the specimen. the specimen, water saturated and placed in a holder, is after a few seconds submitted to a constant flow. At this time the increase of the weight of the out flowing water with time is measured. Based on Darcy's law for laminar flow, the weight per unit time is then recalculated to volume, and the permeability follows from the formula :

$$K = \frac{m \cdot V \cdot L}{A \cdot dP \cdot t}$$

Gas permeameter

The gas permeameter uses pressurized gas to determine the permeability of the specimen in three different ranges of sensitivity. The instrument includes; a thermometer; a flowrate meter; a manometer; and a pressure regulating valve. The kernel is placed in the holder surrounded by a rubber mantle to exclude gas leakage at the sides of the kernel. When gas under 1 atm. pressure is guided through the kernel, the permeability can be calculated according to Darcy from :

$$K = \frac{m \cdot Q \cdot L}{A \cdot P}$$

were m = viscosity (Cp)
 m water = 1 Cp
 m gas = 0.0176 Cp at 23° C
 K = permeability (Darcy)

$$V_2 = \frac{10}{10 - V_1 - V_2} (V_1 - V_2) \quad (2)$$

$$V_2 = 1 - \frac{V_1}{10} \quad (3)$$

Appendix B.3 Permeability measurement

Liquid permeameter

The liquid permeameter is an apparatus which uses pressurized water to create a flow through the specimen. The specimen, water saturated and placed in a holder, is after a few seconds subjected to a constant flow. At this time the increase of the weight of the out flowing water with time is measured. Based on Darcy's law for laminar flow, the weight per unit time is then related to volume, and the permeability follows from the formula:

$$K = \frac{m \cdot V \cdot L}{A \cdot \Delta p \cdot t}$$

Gas permeameter

The gas permeameter uses pressurized gas to determine the permeability of the specimen in three different ranges of sensitivity. The instrument includes: a thermometer; a flowmeter; a sensor; and a pressure regulating valve. The kernel is placed in the holder surrounded by a rubber mantle to exclude gas leakage at the sides of the kernel. When gas under a certain pressure is pushed through the kernel, the permeability can be calculated according to Darcy's law:

$$K = \frac{m \cdot V \cdot L}{A \cdot \Delta p \cdot t}$$

where
 K = permeability (Darcy)
 m = mass (g)
 V = volume (cm³)
 L = thickness (cm)
 A = area (cm²)
 Δp = pressure difference (atm)
 t = time (s)

Appendix A Determination of Parameters

Q = average value of flow rate (cl/sec)
P = pressure difference in atmosphere
L = length of kernel (cm)
A = square area of kernel (cm²)
V = volume of out flow (cc)
t = time (sec)

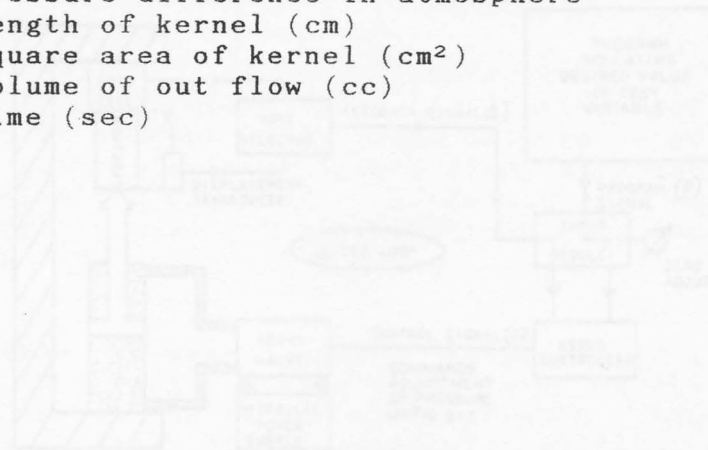


Figure 8.3.1 Closed loop Servo Controlled system

For calibration of the 700 kN loading frame a steel specimen, of elastic modulus 210 GPa, is tested. In figure 8.3.2 the measured value is approximately 200 GPa. In the calibration program of the computer a correction factor for the deformation of the 30 mm of the bottom plate, which is also measured by the LVDT's, is accounted for.



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Figure 8.3.2 Calibration Curve of Steel

Q = average value of flow rate (l/sec)
 P = pressure difference in atmosphere
 L = length of kernel (cm)
 A = square area of kernel (cm²)
 V = volume of oil flow (cc)
 t = time (sec)

Appendix B.3

Unconfined Compressive Strength Determination

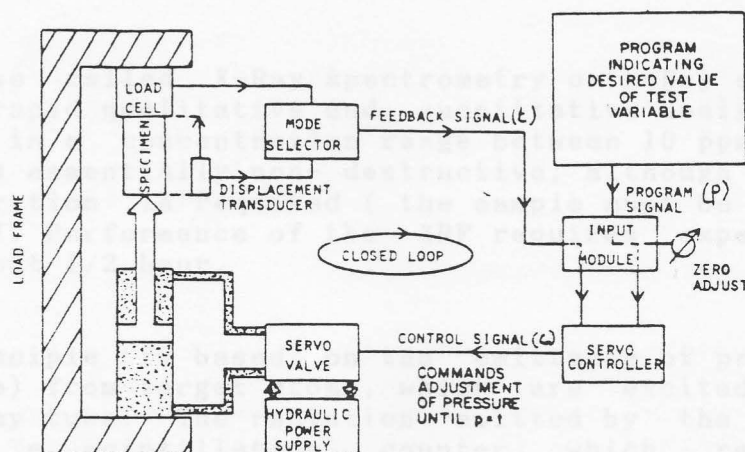
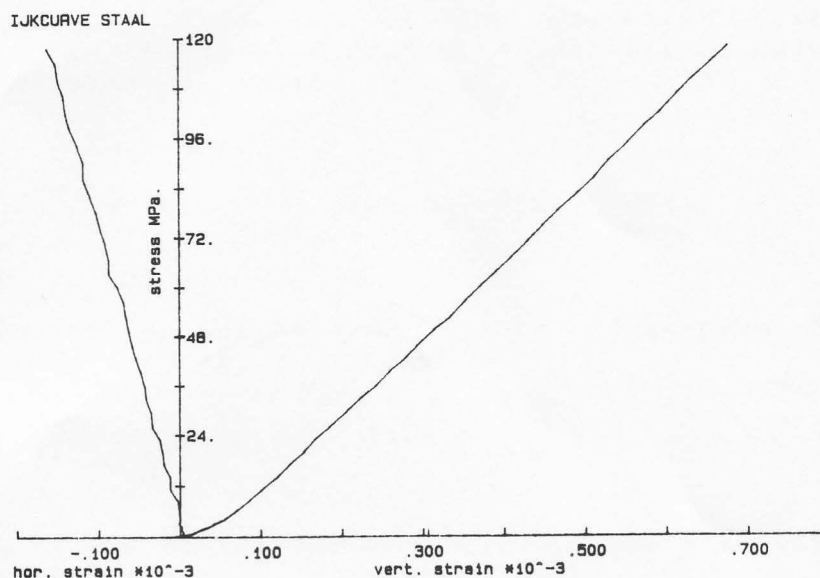


figure B.3.1 Closed Loop Servo Controlled System

For calibration of the 700 kN loading frame a steel core specimen, of elastic modulus 210 GPa, is tested. In figure B.3.1 the measured value is approximately 200 GPa. In the calculation program of the computer a correction factor for the deformation of the 30 mm of the bottom plate, which is also measured by the LVDT's, is accounted for.



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figure B.3.2 Calibration Curve of Steel



Figure B.3.1 Closed Loop Servo Controlled System

For calibration of the 700 kN loading frame a steel core specimen of elastic modulus 210 GPa, is tested. In Figure B.3.1 the measured value is approximately 200 GPa. In the calibration program of the computer a correction factor for the deformation of the 30 mm of the bottom plate, which is also measured by the LVDT's, is accounted for.

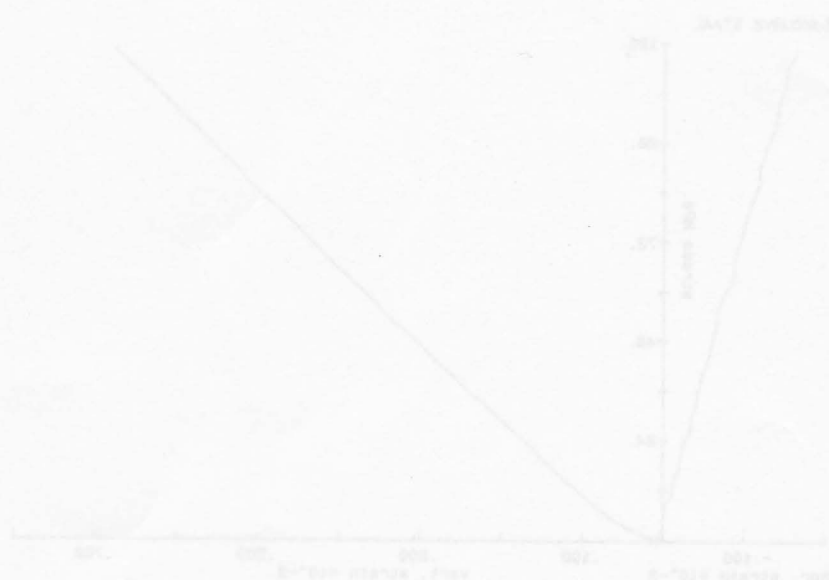


Figure B.3.2 Calibration Curve of Steel

Appendix B.4 X-Ray Fluorescence Analysis

XRF (also called X-Ray spectrometry or X-Ray emission) can be used for rapid qualitative and quantitative analysis of over 80 elements in a concentration range between 10 ppm and 100 %. The method is essentially non-destructive, although some form of sample preparation is required (the sample must be submitted in powder form). Performance of the XRF requires expert knowledge and takes about 1/2 hour.

The principle is based on the emittance of primary X-rays (fluorescence) from target atoms, which are excited by a beam from an X-ray tube. The radiation emitted by the atom is then absorbed by a scintillation counter, which registers the intensities for the different elements. The height and amount of these signals, after amplifying, are then proportional to the quantity and quality of the sample. They can be recorded as the number of pulses per unit time.

$$F(x) = Pr\left\{ \sum_{i=1}^n X_i \leq x \right\} \quad \text{with } n \text{ standard normal}$$

F represents the unknown population under the condition that random sampling set comes from a normal distributed population with μ and σ unknown. A large deviation between F_0 and F will result in the rejection of the suspected normality. As deviation the integrated quadratic difference between F_0 and F will be considered, namely

$$W = n \int_{-\infty}^{\infty} (F_0(x) - F(x))^2 dx$$

This integral can be, because F_0 is a stepped function (C.1.1), redefined as,

$$W = \sum_{i=1}^n \frac{1}{n} (F_0(x_i) - F(x_i))^2 + \left(\frac{1}{n} \right) \sum_{i=1}^n \frac{1}{n} (F_0(x_i) - F(x_i))^2$$

in which $t_i = F(x_i)$ for $i = 1, 2, \dots, n$ and $x(1), x(2), \dots$ arranges the values in increasing height.

The critical value is given by *

XRF (also called X-Ray Spectrometry or X-Ray emission) can be used for rapid qualitative and quantitative analysis of over 50 elements in a concentration range between 10 ppm and 100 % . The method is essentially non-destructive, although some form of sample preparation is required (the sample must be submitted in powder form). Performance of the XRF requires expert knowledge and takes about 1/2 hour.

The principle is based on the emission of primary X-rays (fluorescence) from target atoms, which are excited by a beam from an X-ray tube. The radiation emitted by the atom is then absorbed by a scintillation counter, which registers the intensity for the different elements. The height and amount of these signals after amplifying, are then proportional to the quantity and quality of the sample. They can be recorded as the number of pulses per unit time.

Appendix C.1 Test for Normality

The empirical distribution function analysis is a technique, which does not require very large numbers of data to render information about the normality of a set. Regarding random sampled values, $x_1, x_2, x_3, \dots, x_n$, the empirical distribution function can be determined as:

$$F_n(x) = \frac{k}{n} \quad \text{if } k \text{ measurements } < x$$

This function is nothing more than the frequency quotient of an event " $X < x$ " as a function of x , based on n measured values. Ofcourse the empirical distribution function represents the unknown population distribution. As an approximate of this distribution function the sample set average \bar{x} and the sample set standard deviation s^* can be calculated and the adapted normal distribution function as well.

$$F(x) = \Pr\left\{ u < \frac{x - \bar{x}}{s^*} \right\} \quad \text{with } u \text{ standard normal}$$

F represents the unknown population under the condition that the random sampling set comes from a normal distributed population with m and σ unknown. A large deviation between F_n and F will result in the rejection of the suspected normality. As a deviation the integrated quadratic difference between F_n and F will be considered, namely

$$W = n \int_{-\infty}^{\infty} \{F_n(x) - F(x)\}^2 dF(x)$$

This integral can be, because F_n is a stepped function (fig. C.1.1), redefined as,

$$W = \sum_{i=1}^n t_i - \frac{1}{n} \sum_{i=1}^n (2i-1)t_i + \left(\frac{n}{3}\right) = \frac{1}{12n} + \sum_{i=1}^n \left(t_i - \frac{2i-1}{2n}\right)^2$$

in which $t_i = F(x(i))$ for $i = 1, 2, \dots, n$ and $x(1), x(2), \dots, x(n)$ arranges the values in increasing height.

The critical value is given by $*$,

The empirical distribution function analysis is a technique which does not require very large numbers of data to render information about the normality of a set of measured random sample values, x_1, x_2, \dots, x_n . The empirical distribution function can be determined as:

$$F(x) = \frac{k}{n} \quad \text{if } k \text{ measurements} \leq x$$

This function is nothing more than the frequency quotient of an event " $x \leq x$ " as a function of x , based on a measured value. Of course the empirical distribution function represents the unknown population distribution. As an approximation of this distribution function the sample set average \bar{x} and the sample set standard deviation s can be calculated and the adjusted normal distribution function as well.

$$F(x) = F\left[\frac{x - \bar{x}}{s} \sqrt{n}\right] \quad \text{with a standard normal}$$

F represents the known population under the condition that the random sampling set comes from a normal distributed population with μ and σ unknown. A large deviation between \bar{x} and \bar{y} will result in the rejection of the suspected normality. As a deviation the integrated quadratic difference between \bar{x} and \bar{y} will be considered, namely

$$W = n \int_{-\infty}^{\infty} [F(x) - \bar{F}(x)]^2 dF(x)$$

This integral can be determined because F is a stepped function (Fig. B.1.1), redefined as:

$$F(x) = \frac{k}{n} = \frac{1}{n} \sum_{i=1}^k 1 = \frac{1}{n} \sum_{i=1}^k \delta(x - x_i)$$

in which $\delta = \delta(x - x_i)$ for $i = 1, 2, \dots, n$ and x_1, x_2, \dots, x_n arranges the values in increasing height.

The critical value is given by

$\frac{0.104}{1 + (1/2n)}$	if the unreliability $a = 0.10$
$\frac{0.126}{1 + (1/2n)}$	if the unreliability $a = 0.05$
$\frac{0.178}{1 + (1/2n)}$	if the unreliability $a = 0.01$

*) Biometrika Tables for statisticians, Vol. 2, Cambridge University Press (1972), table 54 .

0.104	at the unreliability $\alpha = 0.10$
$T = (1/2\pi)$	
0.128	at the unreliability $\alpha = 0.05$
$T = (1/2\pi)$	
0.178	at the unreliability $\alpha = 0.01$
$T = (1/2\pi)$	

Biometrika Tables for Statisticians, Vol. 2, Cambridge University Press (1972), Table 64

