Fatigue Behaviour of Steel Reinforcement Bars at Very High Number of Cycles

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“In fatigue and solitude men emanate the divine.”
— Louis-Ferdinand Céline, Journey to the End of the Night
Today’s engineering methods of fatigue safety verification of existing civil structures like reinforced concrete bridges are still based on conservative methodologies used for the design of new structures and on incomplete knowledge on the fatigue behaviour of steel rebars in particular in the relevant domain of very high number of cycles at relatively low fatigue stresses, as experienced by reinforced concrete bridge deck slabs.

This situation often leads to the unsatisfactory situation that the fatigue safety of existing bridges cannot be verified “on paper” although there are no signs of any fatigue damage. This situation may lead to costly unnecessary interventions for strengthening or even replacement of bridges. Consequently, there is an obvious need to improve knowledge about the fatigue behaviour of reinforced concrete elements subjected to very high number of stress cycles, in particular for steel reinforcing bars which are the fatigue vulnerable part of reinforced concrete.

In her doctoral thesis, Marina Rocha investigates the fatigue behaviour of steel reinforcement bars at very high number of stress cycles using concepts of mechanics of materials including micro and macro structural material aspects and fracture mechanics. By an increase in knowledge about the fatigue behaviour of reinforced concrete, more realistic methods for the examination of existing bridges will be developed contributing to devise novel ways how to “get more out of an existing structure subjected to fatigue”. This thesis contributes to this ambitious goal and is thus much relevant from a socio-economic viewpoint and sustainability of civil structures.

The doctoral thesis by Marina Rocha contains a significant amount of new data and information on the fatigue behaviour of steel reinforcement bars subjected to very high number of stress cycles. The research includes fatigue test results at very high number of stress cycles, micro-macro structural material characterisation of
steel rebars as well as numerical and analytical modelling to investigate the main parameters influencing the fatigue behaviour and fatigue strength of steel rebars.

With her doctoral thesis, Marina Rocha provides the proof of her capabilities to conduct a significant scientific study and to solve complex scientific questions by applying scientific approaches. The present thesis delivers results and findings that are useful and applicable to improve structural engineering methods for the fatigue safety verification of existing reinforced concrete structures. In the name of the whole MCS Team, I thank Marina for her constant and thorough investment in the thesis topic as well as for her professional skills and personal qualities.

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Abstract

A large portion of reinforced concrete (RC) bridges in the western world were built in the second half of the last century. RC bridge deck slabs are nowadays often subjected to increased traffic loading and volume than originally designed for and thus, steel reinforcement bars (rebars) are more susceptible to fatigue damage. Fatigue life of rebar can be largely affected by the crack initiation phase characterised by the growth of short cracks. The approaches available for fatigue damage evaluation of rebar fail to predict the crack initiation phase. Microstructural barriers control the short crack behaviour which can be significantly different from the stable long crack growth described by Paris’ law. The stochastic nature of the fatigue life comes mainly from the scatter of these short cracks. Research on this domain is attractive since it can help to understand more accurately the fatigue behaviour of rebar. A better understanding can result in more accurate fatigue damage evaluation of RC elements.

The aim of this thesis is to predict the scatter and fatigue behaviour of hot rolled (HR), cold worked (CW) as well as quenched and self-tempered (QST) rebar incorporating the crack initiation phase. The research commences with an experimental investigation on the fatigue strength of QST rebar under high and very high cycle fatigue (HCF-VHCF) at constant amplitude (R=0.1). A non-destructive inspection technique was applied for surface crack detection based on the frequency change monitored during the tests.

Surface and near surface macro residual stresses on QST rebars were determined by X-ray diffraction and Cut Compliance techniques. Surface imperfections and roughness were identified with Scanning Electron Microscopy mainly near the ribs. A parametric study of the rebar geometry, using 3D Finite Element Models, allowed to determine the influence of rib inclination and rebar diameter on the stress concentration factors.
A short crack growth model was developed to study the scatter resulting from the interaction between short crack and microstructural barriers. The model includes dispersion of grain orientation ratio, grain size variation and different phases (ferrite-pearlite and martensite). This model was then modified to include the surface roughness effects and long crack propagation. The stress concentration factor was considered as a constant parameter. The model predicted the fatigue behaviour of HR-CW and QST rebars.

**Keywords:** Steel reinforcement bars; Fatigue tests under HCF-VHCF; Micro-macro structural surface characterisation; Constant amplitude; Short crack growth model; Scatter; Fatigue behaviour prediction.
Une grande partie des ponts en béton armé (BA) dans le monde occidental ont été construits au cours de la seconde moitié du siècle dernier. Les dalles en BA de ponts sont aujourd’hui soumises à des charges et volumes accrus vis-à-vis de leur conception initiale et par conséquent, les barres d’armature en acier sont plus sensibles aux dommages de fatigue résultants de l’augmentation du trafic. La résistance à la fatigue des armatures peut-être en grande part influencée par la phase d’initiation des fissures — caractérisée par la croissance de fissures courtes. Les approches disponibles pour l’évaluation des dommages en fatigue des armatures ne parviennent pas à prédire la phase d’initiation des fissures. Les barrières microstructurales contrôlent le comportement des fissures courtes qui diffère de celui des fissures longues décrites par la loi de Paris en régime stable. La nature stochastique de la vie en fatigue provient principalement de la dispersion générée par les fissures courtes. La recherche dans ce domaine est intéressante, car elle peut aider à appréhender plus précisément le comportement en fatigue des armatures. Une meilleure compréhension entraînera une plus grande précision de l’évaluation des dommages en fatigue des éléments en BA.

L’objectif de cette thèse est de prédire la dispersion et le comportement des essais de fatigue des barres d’armature laminées à chaud (HR), travaillées à froid (CW) ainsi que des barres trempées et revenus (QST) en considérant la phase d’initiation des fissures. Le projet de recherche débute par une étude expérimentale de la résistance à la fatigue des armatures QST à grand et très grand nombre de cycles (HCF-VHCF) à amplitude constante (R=0.1). Une technique d’inspection non destructive a été appliquée pour la détection des fissures de surface se basant sur la variation de la fréquence mesurée pendant les essais.

Les contraintes résiduelles macroscopiques de surface et près de la surface des armatures QST ont été déterminées par diffraction des rayons X et par la méthode
"Cut Compliance". Les imperfections de surface et la rugosité ont été identifiées principalement près des nervures par microscopie électronique à balayage. Une étude paramétrique de la géométrie des barres d’armature, à l’aide de modèles 3D par éléments finis, a permis de déterminer l’influence de l’inclinaison de la nervure et du diamètre des barres d’armature sur les facteurs de concentration des contraintes.

Un modèle de propagation des fissures courtes a été développé pour étudier la dispersion résultant de l’interaction entre les fissures courtes et les barrières microstructuralles. Le modèle inclut la dispersion liée à l’orientation cristallographique des grains, à la variation de la taille des grains et des phases (ferrite-perlute et martensite). Ce modèle a ensuite été amélioré pour inclure les effets de la rugosité de surface ainsi que la phase de propagation linéaire des fissures longues. Le facteur de concentration de contrainte a été considéré comme un paramètre constant. Le modèle a prédit le comportement en fatigue des armatures HR-CW et QST.

**Mots-clés**: Barres d’armature en acier ; Essais en fatigue HCF-VHCF ; Caractérisation microstructurale et macrostructurale de surface ; Modèle de propagation de fissures courtes ; Dispersion ; Prévision du comportement en fatigue.
Resumo

Uma grande parte das pontes de concreto armado (CA) no mundo ocidental foi construída na segunda metade do século passado. Atualmente, as lajes em CA das pontes estão sujeitas a um aumento de carga e volume de tráfego em relação ao projeto inicial e, consequentemente, as barras de aço estão mais suscetíveis a danos por fadiga. A vida à fadiga das barras pode ser, em grande parte, influenciada pela fase de iniciação das fissuras caracterizada pelo crescimento de microfissuras. Os métodos utilizados para avaliação dos danos por fadiga em barras de aço são incapazes de prever essa fase de iniciação. Barreiras microestruturais controlam o comportamento das microfissuras o qual pode diferenciar significativamente do crescimento estável das fissuras descrito pela lei de Paris. A natureza estocática da vida à fadiga é principalmente influenciada pela dispersão gerada pelas microfissuras. Pesquisa nessa área é relevante pois contribui para um melhor entendimento do comportamento à fadiga das barras e, portanto, uma avaliação mais precisa do dano por fadiga em elementos de CA.

O objetivo desta tese é prever a dispersão e comportamento à fadiga das barras de aço laminadas a quente (HR), trabalhadas a frio (CW) assim como temperadas e revenidas (QST) considerando a fase de iniciação das fissuras. Este projeto de pesquisa inicia-se por uma investigação experimental da resistência à fadiga das barras QST sujeitas a alto e muito alto ciclos à amplitude constante (R=0.1). Uma técnica de inspeção não destrutiva foi utilizada para identificação de fissuras na superfície da barra baseada na variação da frequência monitorada durante os ensaios.

Tensões residuais macroscópicas foram determinadas na superfície e subsuperfície das barras QST por difração de raio X e o método “Cut Compliance”. Impefeições e rugosidade da superfície foram identificadas principalmente próximas aos dentes das barras pelo microscópio eletrônico de varredura. Um estudo paramétrico da geometria das barras usando modelos 3D de elementos finitos permitiu deter-
minar a influência da inclinação dos dentes e do diâmetro das barras nos fatores de concentração de tensão.

Um modelo de crescimento de microfissuras foi desenvolvido para estudo da dispersão resultante da interação entre fissuras e barreiras microestruturais. O modelo inclui a dispersão da razão de orientação dos grãos, variação do tamanho do grão e diferentes fases (ferrita-perlita e martensita). Este modelo foi posteriormente modificado para inclusão dos efeitos da rugosidade da superfície e propagação de macrofissuras. O fator de concentração de tensão foi considerado como um parâmetro constante. O modelo previu o comportamento à fatiga das barras HR, CW e QST.

**Palavras-chave:** Barras de aço; Ensaios de fatiga a HCF-VHCF; Caracterização micro e macroestrutural da superfície; Amplitude constante; Modelo de crescimento das microfissuras; Dispersão; Previsão do comportamento à fatiga.
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Chapter 1

Introduction

Context and motivation

In the second half of the last century, a significant increase in bridge construction was observed. This event was accompanied by the change from steel to reinforced concrete (RC) as the dominant structural material [1].

A survey performed on approximately 50 000 concrete bridges, (RC, prestressed and post-tensioned), by the railway administration in 17 European countries (including Switzerland), showed that about 80% of these bridges are classified as RC. From all concrete bridge types, 25% are less than 20 years old, 55% are between 20 and 50 years old, 16% are between 50 and 100 years old and 4% are over 100 years old [2].

In the United States, bridges have been mostly built in the last 50 years with RC material being predominant. RC bridges correspond to approximately 30% (or more than 140 000) of all bridge types in the United States [3].

Bridges are nowadays often subjected to increased traffic loading and volumes than originally designed for. RC bridge deck slabs can experience very high number of significant stress cycles during their service life i.e., exceeding 10 million cycles [3]. Thus, they are more susceptible to fatigue damage with a likely ultimate strength reduction of the building materials due to fatigue. Despite this fact, reinforced concrete deck slabs were commonly not designed for fatigue [4]. RC bridges built before 1989 in Switzerland, for example, often do not meet the current fatigue code requirements [5].

[4] and [6] showed that fatigue failure in slab-like members is always induced by the fracture of steel reinforcement bars (rebars). They are significantly more fatigue vulnerable than concrete which shows no or minor local fatigue damaging [4]. Therefore, the knowledge on the fatigue behaviour of rebars is of fundamental im-
portance for safety evaluation of RC elements.

Research on the fatigue strength of rebars was intensely carried out up to the 1980's on hot rolled (HR) and cold worked (CW) steels [7], [8] and [9]. Axial and bending fatigue tests were performed, under constant stress amplitude, on HR and CW rebars in air and embedded in concrete, respectively. These test results formed the basis of the standard S-N curves used still nowadays for fatigue safety verification. In the last decades, straigth HR and CW steels have been replaced by quenched and self-tempered (QST) rebars. In some European countries, QST rebars were introduced in 1974 [10]. Straigth QST rebars correspond to approximately 1/3 of the Swiss rebar market. Rebars with diameter greater than 20 mm are mainly QST steels; rebars with diameter smaller than 20 mm are mainly cold worked steels with very small diameters consisting of smooth cold worked bars. QST rebars are hot rolled steels followed by rapid water quenching. This process results in a harder outer surface layer compared to HR and CW rebars.

Although a large amount of experimental data of QST rebars is available, the test results are often limited to 2 million stress cycles with rarely tests exceeding 5 million cycles. These rebars have mostly shown improved fatigue strength and smaller scatter in the tests compared to HR and CW rebars.

The fatigue damage in rebars develops in two stages: crack initiation with nucleation and growth of short (micro) cracks followed by a linear long (macro) crack propagation until failure occurs. Rebar surface is a preferential site for fatigue crack initiation. Surface conditions may significantly influence the behaviour of short cracks. The understanding of the mechanisms of crack initiation is therefore a key issue.

In a damage-tolerance approach, it is assumed that a rebar contains an initial long crack. This crack propagates according to the Paris' law and since failure of one rebar is not failure of the slab, a different resistance factor is used. However, short cracks can behave significantly different from the long crack propagation predicted by Paris' law [11]. In addition, S-N curve approach used for fatigue life assessment, although relatively simple and widely used, doesn't separate crack initiation and propagation phases.

Short crack growth models have been widely used to predict the fatigue behaviour of metals. The significance of microstructural features on the fatigue damage process is considered by these models. They can predict the behaviour of short cracks by simulating the interactions between crack and microstructural barriers.
1.1 Objectives of thesis

The main objectives of this thesis are summarised as follows:

1. Investigate experimentally the fatigue strength of QST rebars at high and very high number of constant amplitude stress cycles.
2. Characterise micro-macro structural aspects of QST rebars.
3. Investigate the influence of surface microstructure on the scatter above the fatigue limit of HR-CW and QST rebars.
4. Predict the scatter and fatigue behaviour of HR-CW and QST rebars.

1.2 Scope of thesis

This thesis focuses on the influence of surface microstructural features on the fatigue behaviour of rebars in the high and very high cycle domain. Both initiation (nucleation and short crack growth) and propagation of long fatigue cracks are important in understanding their behaviour. However, the prime importance of the crack initiation phase is addressed in this thesis.

In this research, microcracks or cracks coalescence are not considered. The analyses are restricted to fatigue under constant amplitude loadings (sequence, interaction effects are not studied). This research includes only straight rebars without concrete; no welding (rebars meshes) are analysed. The study covers only rebars tested under positive R-ratio (R=0 to 0.2). The QST rebars analysed in this thesis were produced by Stahl Gerlafingen AG.

Fatigue tests with QST rebars carried out at high and very high number of stress cycles are discussed. It is followed by an experimental investigation of micro-macro structural surface conditions on QST rebars. A parametric study allowed to determine the influence of rib geometry on the stress concentration factors at the rebar surface. A model was then developed to quantify the contribution of microstructural parameters on the scatter found in fatigue tests. Although the experimental analyses were performed only on QST rebars, the model was also adapted to study the scatter on HR-CW rebars, present in older RC bridges. This model simulates the interactions between short crack and microstructural barriers. It considers stochastic crack initiation and long crack propagation phases. The model was applied for fatigue behaviour prediction of HR-CW and QST rebars and to investigate the scatter in fatigue tests. The model results were compared to experimental data.
1.3 Structure of thesis

The structure of this thesis and its contents are presented in Fig. 1.1. The thesis is divided in three parts: 1) Experimental, with fatigue test results and rebar characterisation 2) Theoretical, where a short crack model is developed 3) Application of the model to the scatter and fatigue behaviour prediction on experimental data.

This research consists of four journal papers, to be submitted, and appendices presented in an extended format.

Chapter 2 presents axial fatigue test results of QST rebars between $10^6$ and $10^8$ stress cycles under constant amplitude, $R=0.1$. Gripping methods are investigated and a non-destructive inspection technique is proposed to detect surface cracks.

Chapter 3 provides experimental investigation on surface and near surface residual stresses and surface imperfections on QST rebars. A parametric study using 3D Finite Element models is developed to determine the stress concentrations factors on the rebar surface.

Chapter 4 presents a short crack growth model, adapted from Navarro and De Los Rios [12], which considers the dispersion in the grain orientation ratio, grain size variation and different phases on the scatter observed in experimental data as obtained above the fatigue limit.

Chapter 5 presents modifications on the model developed in Chapter 4 to include surface roughness dispersion and long crack propagation. The stress concentration factor from the rib geometry is considered in the calculations. The model results are compared to experimental data.

Finally, Chapter 6 summarises the main conclusions of the four previous or main chapters and suggests areas for future work in this field.

Supplementary information are presented in Appendices A to D which consist of:

- 3D Finite Element Models of the rebar geometry and stress concentrations analyses for different rib geometries;
- the algorithm developed for the short crack growth model presented in Chapters 3 and 4;
- the 3D roughness profile determined from the surface of the QST rebar;
- a Conference paper where an approach was proposed to determine S-N curve from fatigue test results including run-out results.
Figure 1.1 – Structure of the thesis.
Bibliography


Very high cycle fatigue tests of quenched and self-tempered steel reinforcement bars
— Marina Rocha, Silvain Michel, Eugen Brühwiler, Alain Nussbaumer

Abstract: Investigations on the fatigue strength of steel reinforcement bars (rebars) mainly involves fatigue tests with hot rolled (HR) and and cold worked (CW) steels. However, in the last few decades, HR and CW rebars were replaced by quenched and self-tempered (QST) rebars with hardened surface layer. There still remains a lack of research on fatigue strength of QST rebars especially in the very high cycle domain i.e., number of stress cycles surpassing 5 million. This work aims to investigate the fatigue performance of QST rebars axially tested at number of stress cycles in the range of $10^6$ to $10^8$. A preliminary study of the gripping method is followed by fatigue test results including non-destructive inspection of the rebar surface and fractographic analyses. The rebar surface is examined with liquid penetrant to reveal fatigue crack location and size in specific frequency interval monitored during the tests. Fractured surface analyses are performed by Scanning Electron Microscopy (SEM) to detect the location from where fatigue cracks initiate. Cross sectional area reduction resulting from fatigue crack propagation is also determined. Fractographic investigations are compared with the fractured surfaces of HR, CW and QST rebars from the literature.

Keywords: Quenched and self-tempered rebars; High and very high cycle fatigue; Gripping method; Non-destructive inspection; Fractured surface analysis.
2.1 Introduction

Reinforced concrete structures such as bridges are nowadays subjected to higher and more frequent traffic loads and thus they are more susceptible to fatigue damage. One of the key elements contributing to the bridge deck slab service life is the fatigue strength of steel reinforcement bars (rebars). Fatigue loading may lead to failure of rebars in the reinforced concrete without any sign of external structural distress except local concrete cracking.

Axial and bending tests of plain rebars and within concrete beams respectively are the two test methods commonly used to study the fatigue strength of rebars. Generally, fatigue tests on rebars are carried out as repetitive loading with stress ratio between 0 and 0.2 [1–3]. Axial fatigue tests on rebars are usually conducted on electromagnetic resonance machines at frequencies up to 150 Hz [4]. The disadvantage of these tests is related to the method of gripping the rebar. It tends to cause local stress concentration and premature failure of the rebar in the gripping area which are not characteristic of the rebar itself. Bending fatigue tests have the advantage of simulating the service conditions at the steel-concrete interface. However, concrete beams are usually tested by hydraulic machines at frequency smaller than 10 Hz, with few tests conducted for number of cycles more than $10^7$, due to the high costs [1].
Fatigue tests carried out up to the 1980’s were mainly on hot rolled (HR) and cold worked (CW) rebars. However, HR and CW rebars were replaced in most European countries by quenched and self-tempered (QST) rebars [5, 6]. These rebars have a hard outer layer of martensite as a result of the specific QST treatment. This process known as Thermex or Tempcore has been introduced in Western Europe since 1974 [7].

Axial and bending fatigue tests performed on QST rebars have been reported in the literature. Thandavamoorthy [8] conducted fatigue tests with Tempcore rebars in eight concrete beams up to 2 million cycles; fatigue strength of QST rebars was found to be comparable to HR and CW rebars. In [3], Tempcore rebars survived to stress levels as high as 40% of the tensile strength \( \sigma_u \) and in some cases reached 60% up to 2 million cycles. Surface imperfections and stress concentrations arising from the rib geometry were significant factors affecting the fatigue lifetime of the fractured rebars. Axial fatigue tests with Tempcore rebars were performed to a maximum of 5 million cycles [9]. The test results showed small scatter for rebars with different diameters. In [10], fatigue tests with HR, CW and Tempcore rebars were run to utmost 2 million cycles. Tempcore rebars showed considerably smaller scatter and higher fatigue strength than HR and CW rebars. In [11], concrete beams with embedded 12 mm diameter Thermex rebars were tested to utmost 10 million cycles. Rebars survived at stress levels higher than 40% of their yield strength \( \sigma_y \).

Fatigue tests performed on QST rebars are mostly limited to utmost 5 million cycles. Thus, fatigue resistance of QST rebars based on these test data can lead to incoherent resistance estimation in the very high cycle regime [12] i.e., beyond 5 million cycles. This paper presents an experimental investigation carried out on QST rebars in the very high cycle fatigue regime. The gripping arrangement used in the axial fatigue tests are discussed. The test frequency is monitored for fatigue crack detection using liquid penetrant testing. The fractured surfaces are analysed by Scanning Electron Microscopy (SEM) and sites where fatigue cracks initiate are identified. Crack propagation region is estimated after test stopping.

### 2.2 Material properties

The chemical composition and mechanical properties of QST (Thermex) rebars with diameter of 16 mm were provided by the manufacturer and are summarised in Tables 2.1 and 2.2.
Table 2.1 – Chemical composition of the QST rebar with diameter of 16 mm.

<table>
<thead>
<tr>
<th>Elements</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>C</td>
<td>0.186</td>
</tr>
<tr>
<td>Si</td>
<td>0.22</td>
</tr>
<tr>
<td>Mn</td>
<td>0.86</td>
</tr>
<tr>
<td>P</td>
<td>0.023</td>
</tr>
<tr>
<td>S</td>
<td>0.040</td>
</tr>
<tr>
<td>Cr</td>
<td>0.14</td>
</tr>
<tr>
<td>Mo</td>
<td>0.02</td>
</tr>
<tr>
<td>Ni</td>
<td>0.13</td>
</tr>
<tr>
<td>Cu</td>
<td>0.40</td>
</tr>
<tr>
<td>Sn</td>
<td>0.018</td>
</tr>
<tr>
<td>V</td>
<td>0.002</td>
</tr>
<tr>
<td>Nb</td>
<td>0.002</td>
</tr>
<tr>
<td>Al</td>
<td>0.004</td>
</tr>
<tr>
<td>Ceq*</td>
<td>0.398</td>
</tr>
</tbody>
</table>

Notes:

(*) Ceq: Carbon Equivalent Ceq (%) = C (%) + $\frac{Mn}{6}$ (%) + $\frac{Cr+Mo+V}{5}$ (%) + $\frac{Ni+Cu}{15}$ (%)

Table 2.2 – Mechanical properties of the QST rebar with diameter of 16 mm.

<table>
<thead>
<tr>
<th>$\sigma_y$ (MPa)</th>
<th>$\sigma_u$ (MPa)</th>
<th>$\varepsilon_{u}$ (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>M</td>
<td>SD</td>
<td>M</td>
</tr>
<tr>
<td>518</td>
<td>8.29</td>
<td>613</td>
</tr>
</tbody>
</table>

Notes:

$\sigma_y$: yield strength;
$\sigma_u$: tensile strength;
$\varepsilon_{u}$: strain at maximum force;
M: mean;
SD: standard deviation.
The microstructure consists of a hardened outer layer of martensite as indicated in Fig. 2.1 with a thickness of approximately 1.2 mm and a soft core of ferrite-pearlite. Vicker’s hardness, measured in the cross section, varied from 265 HV in the martensitic layer to 160 HV in the core; these values are expected for QST rebars as given in [3, 6].

Figure 2.1 – Etched cross section of a 16 mm diameter QST rebar.

2.3 Test details

2.3.1 Grip arrangement and specimen preparation.

Axial fatigue tests are sensitive to the high stress concentration induced by the gripping pressure preventing the rebar from slipping. Some techniques were investigated to avoid premature failure of the rebar in the grip area.

The initial grip system used in the testing machine is shown in Fig. 2.2a. The grip has an inner-circular cross section of 20 mm. Three types of gripping arrangement were used with the initial system in order to obtain the failure in the rebar free length: 1) A 1 mm aluminium sheet was wrapped around the rebar ends within the grip area with the aim to distribute the force evenly over the surface of the rebar as shown in Fig. 2.2b; 2) Shot peening the rebar ends (see Fig. 2.2c) to induce compressive residual stresses on the surface and 3) Welding and machining the rebar ends to create a gradual and smooth transition between the grips and the ribbed surface Fig. 2.2d. The cross section diameter of the welded and machined rebar ends was 20 mm. However, all these methods were ineffective to prevent failure in the grip area.

Therefore, the initial grip system was replaced by a conical grip with maximum 18 mm diameter (see Figs. 2.3a and 2.3b) in the testing machine. Different conical
Figure 2.2 – (a) Initial grip system used for the fatigue tests. Rebar failure in the grip area with (b) Aluminium sheet; (c) Shot peened rebar ends; (d) Welded and machined rebar.

arrangements have been shown to be effective for fatigue tests with rebars. However it required casting the rebar ends in alloys as given in [3,9]. In [7], QST rebar ends embedded in high strength metallic grout within a conical grip system showed 40% of the failures in the grips and 60% within a distance of 1 diameter outside the grip area.

In this work, three specimens were initially tested with the conical gripping system to verify its effectiveness. Failures occurred at least 39 mm away from the grip edges without modifying the rebar ends.

The specimen preparation was as follows: 16 mm raw material was first examined for defects, scratches and manufacturer’s identification marks. Then it was cut into pieces of 400 mm length. 80 mm on each side were necessary for clamping; therefore the free length was 240 mm. The specimen free length considered in tests
is in accordance with standard recommended procedures [13] where the rebar’s free length in axial tests should be at least 140 mm or 14 times the specimen diameter, whichever is greater. Care was taken to ensure that the free length was free of manufacturer’s identification marks. The rib patterns on both rebar sides are shown in Figs. 2.4a and 2.4b.

### 2.3.2 Test method

Axial fatigue tests were performed on a RUMUL Testonic 100 kN 8601 resonance machine at 85 Hz and force-ratio of 0.1. A total of 21 specimens were tested; 6 tested specimens were considered as non-valid results since the failure occurred in the grip area. The valid tests i.e, rebar with failure in the free length were run to at least $1.89 \times 10^6$ cycles but not longer than $66.2 \times 10^6$ cycles as given in Table 2.3. The failure within the rebar’s free length occurred in more than 70% of the tested
rebars. Whenever the specimen survived the test, it was termed “run-out”. Two run-outs were retested under higher forces. It was assumed fatigue damage in the tested rebars for a surface crack length of at least 5 mm. This is the minimum crack length that can be detected by the non-destructive inspection technique used in this work.

During all the tests that were carried out, the resonance frequency was monitored. The frequency change was used as an indication of cracking in the specimen. A typical evolution of the frequency drop obtained during the fatigue tests is shown in Fig. 2.5. The frequency drop versus number of cycles can be separated in three stages. In stage I, the frequency drops continuously from the beginning of the tests up to approximately $10^6$ cycles. The frequency drop in the beginning of the test was probably caused by the settlement of the specimen in the clamping area. A stabilization of the frequency is observed at stage II with the frequency variation being smaller than 0.1%. An abrupt frequency change occurs at stage III caused by fatigue crack propagation followed by failure of the specimen.

The test was interrupted just after the abrupt frequency change for non-destructive inspection by liquid penetrant for detection of fatigue cracks. The specimen remained mounted in the machine and loaded at the mean force while the liquid penetrant was applied on the surface. The liquid penetrant was applied only once. The test was stopped at a frequency drop of approximately 2.2%.

![Figure 2.5](image)

**Figure 2.5** – Representative frequency drop (%) of a failed specimen during the fatigue test.
Table 2.3 – Fatigue test results of 16 mm rebar.

<table>
<thead>
<tr>
<th>Specimen number</th>
<th>Stress range (MPa)</th>
<th>Number of cycles ($\times 10^6$)</th>
<th>Failure x Run-out o</th>
<th>Failure location$^{(1)}$ (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>225</td>
<td>30</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>2</td>
<td>235</td>
<td>30</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>235</td>
<td>35</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>4</td>
<td>243</td>
<td>36</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>5</td>
<td>243</td>
<td>51</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>6</td>
<td>243</td>
<td>49</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>7</td>
<td>245</td>
<td>1.89</td>
<td>X</td>
<td>92</td>
</tr>
<tr>
<td>8</td>
<td>247</td>
<td>30</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>9</td>
<td>247</td>
<td>30</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>10</td>
<td>247</td>
<td>30</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>11$^{(2)}$</td>
<td>251</td>
<td>64.5</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>12</td>
<td>251</td>
<td>66.2</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>13$^{(2)}$</td>
<td>255</td>
<td>34.6</td>
<td>O</td>
<td>-</td>
</tr>
<tr>
<td>14</td>
<td>255</td>
<td>2.71</td>
<td>X</td>
<td>60</td>
</tr>
<tr>
<td>15</td>
<td>255</td>
<td>1.97</td>
<td>X</td>
<td>39</td>
</tr>
</tbody>
</table>

Notes:

$^{(1)}$ Failure location: distance from the grip’s edge to the failure in the free length.

$^{(2)}$ Specimens 11 and 13 correspond to the retested 8 and 9 specimens at 251 MPa and 255 MPa respectively.

2.4 Results and discussions

2.4.1 Test results

Table 2.3 shows the fatigue test results of the 16 mm rebars. Two specimens were retested at higher stresses after run-outs. Three rebars showed a failure and all the others were run-outs. The failure location on the rebars are given in Table 2.3. Run-outs represented 80% of the test results with one rebar surviving to a total of 94.5 $\times 10^6$ cycles for a stress level of approximately 48% of the mean yield strength $\sigma_y$.

Fatigue strength of rebars are traditionally expressed by S-N curves. S-N curves are obtained by linear regression applied only to failed data points; run-outs are neglected in the analysis. In the standard S-N curves, a minimum of 12 specimens is required for characteristic allowable and reliability data as given in [14]. The 3 failed specimens in this present work don’t fit the minimum specimen size requirements for linear regression analysis. Therefore, the S-N curve for the present test results couldn’t be determined. In [12], an alternative approach is proposed for statistical treatment of the data including run-out specimens.
2.4.2 Non-destructive inspection

Non-destructive inspection allowed identifying the fatigue crack location and size on the surface of the failed specimens. The liquid penetrant was applied at the beginning of stage III as indicated in Fig. 2.5.

The frequency change measured from the beginning of stage I until application of the liquid penetrant on the 3 failed specimens varied between 0.15 and 0.18%. Fig. 2.6 shows the frequency evolution obtained for the failed specimens 7, 14 and 15. Two dots of penetrant ink shown in Fig. 2.7a indicated the surface crack tips detected just after the abrupt frequency change in specimen 15. The distance between the crack tips was approximately 8 mm. Similar crack length was also detected on the surface of specimens 7 and 14. The crack propagated away from the non-uniform ribs and perpendicular to the longitudinal specimen axis. Fig. 2.7b shows the crack after the test stopping when the frequency dropped by 2.2%. The crack area was approximately 50% of the specimen cross section.

Figure 2.6 – Frequency drop versus number of cycles obtained for specimens 7, 14 & 15.

The frequency drop versus number of cycles obtained for run-out specimens is given in Figs. 2.8 to 2.11. The frequency evolution of run-out specimens showed a similar tendency for stages I and II: A continuous frequency drop followed by an stabilization period until test stopping. The liquid penetrant was applied on the run-outs mounted in the machine at the end of the tests. However, no crack was detected
Figure 2.7 – (a) Specimen 15: Two dots of penetrant ink indicating the fatigue crack tips on the specimen surface; (b) Crack on the specimen surface when the test was stopped.

on the surface of any surviving specimen, even for specimen 13 which showed a frequency drop of nearly 0.5%.

Figure 2.8 – Frequency drop versus number of cycles obtained for specimens 1, 2 & 3.
2.4.3 Fractured surface analyses

Fractography analysis was performed by Optical Microscopy (OM) and XL30-FEG SEM in order to determine the location where fatigue cracks initiate. Since the tests were stopped before complete fracture of the specimens, it was required to split them and prepare the fractured surfaces before microscopic analysis. After test stopping,
the specimen was put in liquid nitrogen in order to split them in a brittle manner using an actuator. The average temperature measured on the specimen surfaces was approximately -65°C after 30 min immersed in liquid nitrogen. QST rebars tends to have a brittle fracture at this temperature [15]. The frozen specimen surfaces were dried using ethanol and compressed air and then left in desiccator under vacuum for one day. The fractured surfaces were then immersed in a beaker containing melted paraffin at 55°C to protect them from damage. A cut was made at approximately 5 cm away from the fractured surfaces by Electric Discharge Machining.

The paraffin around the specimen’s cross section was manually removed. The cross sections were then immersed in an ultrasonic cleaner containing Xylene for 10 minutes in order to remove the residual paraffin. Since some corrosion was visible on the fractured surfaces, two methods of removing corrosion were used: 1) One side of the cross section was immersed in a beaker with Alconox solution heated up to 90°C for 1 hour and 2) The other side was immersed in an ultrasonic cleaner for 10s containing an acid solution, consisting of 3 mL of hydrochloric acid, 4 mL of 2-butyne-1, 4-diol (35% aqueous solution) and 50 mL of deionized water [16]. Both methods were effective to remove the corrosion on the surface. After corrosion removal, the fractured surfaces were cleaned with ethanol, dried with compressed air and left in desiccator under vacuum to protect it from any damage before microscopic analysis.
The site from where fatigue cracks initiate on the specimen surface is indicated in Figs. 2.12a, 2.13a and 2.14a. The crack initiated at or very near the base of the transverse non-uniform ribs. Figs. 2.12b, 2.13b and 2.14b show the fatigue crack propagation region determined from fractography images obtained with OM. The bright rough area is the brittle fracture caused by the actuator whereas the smooth area is the fatigue crack region. The different surface texture allows determining the final fatigue crack area.

Imperfections on the fractured cross section from where fatigue cracks initiated are indicated in Figs. 2.12c, 2.13c and 2.14c. The cracks (white lines in the SEM images) emerging from the imperfections confirm the crack initiation site previously identified in the OM images. A single fatigue crack initiation site is identified on the cross sections of specimens 7 and 14: Cracks start to propagate from a imper-
Figure 2.13 – (a) Location where fatigue crack initiates on the surface of specimen 14; (b) OM image (10×) of the fractured cross section; (c) SEM image (65×) of the imperfection from where fatigue crack initiated.

Fractured surface analyses on HR, CW and QST rebars tested at high number of fatigue cycles have been reported in the literature [3, 4, 17]. According to [4], fatigue lifetime of HR and CW rebars axially tested at high number of cycles was mainly affected by surface defects ranging from 5 to 100 µm. HR rebars mostly had a single fatigue crack initiation site and plane fractured surface. CW rebars had multiple initiation sites and helical fractured surfaces.

In [3], fatigue cracks on QST rebars initiated from surface defects and at the root of the transverse ribs from where arise the highest stress concentration [18].
The plane fractured surface showed single or multiple initiation sites: Higher stress range and lower ratio between rib radius and rib height $r/h$ led to multiple initiation sites.

In [17], fatigue tests performed on HR rebars embedded in concrete beams resulted in fatigue cracks starting at the base of transverse ribs and plane fractured surface. The fractured surface of CW rebars embedded in concrete was plane inclined at an angle of approximately 45° and fatigue cracks initiated near to the transversal ribs.

Comparisons between the fractured surface investigations given in the literature [3, 4, 17] and the fractured surfaces analysed in this present work showed that fatigue life of HR, CW and QST was significantly affected by surface imperfections in axial fatigue tests at high cycle regime. In bending fatigue tests, the rib geometry had a
significant effect on the fatigue lifetime of rebars. Beside, HR and QST rebars tested at high cycle fatigue tended to have a single crack initiation site and a similar plane fractured surface while CW rebars show a helical fractured surface.

2.5 Conclusions

Fatigue tests were performed on QST rebars between \(10^6\) and \(10^8\) cycles and under constant amplitude loading. Non-destructive inspection using liquid penetrant allowed to determine the surface crack size and location just after the abrupt drop of the frequency. Fractured surfaces were analysed after test stopping by OM and SEM and compared to fractographic analysis from the literature. The following conclusions can be drawn from the present study:

- Conical grip arrangement was the only effective method to prevent failure in the grip area. The method provided more than 70% of the failures on rebar free length without requiring any modification at the rebar ends.

- QST rebars survived at least 30 million cycles in 80% of the tests and at stress levels of approximately 50% of the mean yield strength.

- Due to the small frequency change at almost the entire fatigue life of the rebars and the limitation of the penetrant liquid testing in detect surface cracks from few mm, fatigue cracks could only be detected when the rebar approached fracture.

- The fatigue lifetime of QST rebars was significantly controlled by manufacturing imperfections extending from surface to the depth cross section; fatigue cracks initiated from imperfections located at and very near the base of the transversal ribs.

2.6 Acknowledgements

The authors are grateful to Danièle Laub from the Interdisciplinary Centre For Electron Microscopy (CIME) at EPFL for her advices and help with the sample preparation for the microscopic analyses. We are also grateful to Prof. Francesco Stellacci from the Supramolecular Nanomaterials and Interfaces Laboratory (SuNMIL) who provided the laboratory space for the chemical attack and preparation of the samples.
Bibliography


Material and geometrical characterisation of quenched and self-tempered steel reinforcement bars
— Marina Rocha, Eugen Brühwiler, Alain Nussbaumer

Abstract: Quenched and self-tempered (QST) steel reinforcement bar (rebar) is manufactured by Thermex or Tempcore process. Characterisation studies of QST rebars are mostly limited to reveal the hardened outer layer and some improved mechanical properties compared to hot rolled (HR) and cold worked (CW) rebars. However, investigations on residual stresses and imperfections originated from the manufacturing process as well as stress concentrations arising from the ribbed profile are rarely found in the literature. Surface residual stress may be beneficial or detrimental to the fatigue performance of rebars although they have been studied only on the subsurface of QST rebars. Surface imperfections are zones of stress concentration from where fatigue cracks may initiate. Imperfections are usually identified in the fractured cross section resultant from fatigue tests of rebars. Stress concentrations can also arise from the rib geometry. While the rib geometric parameters affect the stress concentration factor $K_t$ on rebars, stress concentration analysis are restricted to 2D Finite Element Models (FEMs). In this present work, characterisation analyses of QST rebars include: 1) Experimental investigation of surface and subsurface residual stresses on QST rebars by Cut Compliance and X-ray diffraction techniques. Residual stresses from both techniques are discussed. 2) Identification of surface imperfections by Scanning Electron Microscopy (SEM) analysis. 3) 3D Finite Element Analysis of stress concentrations on the ribbed profile. The influence of the rib geometry such as radius, width, height and inclination as well as the rebar diameter on $K_t$ values are analysed. The critical zones are determined along the ribs.

Keywords: Quenched and self-tempered rebars; Microstructure; Residual stress; Surface imperfection; Stress concentration factors.
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</tbody>
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### 3.1 Introduction

Concrete structures show typically three types of steel reinforcement bars (rebars): hot rolled (HR), cold worked (CW) and quenched and self-tempered (QST) steels. HR rebars are produced through hot rolling followed by a slow cooling. Billets heated up to about 1200°C pass through rollers which squeeze them into the required diameter. The bars are then air cooled to ambient temperatures after emerge from the last rolling mill with a temperature of about 1000°C. As the rolling finish is above 900°C before the steel is allowed to naturally cool, the resulting properties are similar to those obtained by normalisation process [1]. CW rebars are submitted to strain hardening after hot rolling; cold twisted deformed rebars are produced by stretching and twisting of mild steel, beyond the elastic limit and subsequently releasing of the load [2].
HR and CW rebars were replaced by QST rebars in some European countries [3–5] i.e., hot rolled steels followed by a special thermal treatment. This thermal process known as Thermex is similar to the Tempcore process developed in 1974 [6]. QST rebars show advantages with respect to weldability, bendability and ductility compared to HR and CW rebars [4, 5, 7, 8].

Thermex process combines the conventional hot rolling with rapid water quenching which results in a hardened outer layer of martensite; HR and CW rebars show ductile ferritic-pearlitic microstructure [9]. The biggest temperature gradients are produced in the quenching stage and hence the greatest residual stresses. As freshly quenched steel tend to crack, the tempering process starts just after quenching to minimize distortion, cracking and residual stresses [10].

The residual stresses pattern in QST steels are mainly influenced by the combination of thermal and microstructural changes in the cross section during the fabrication process [10]. Residual stresses on rebars act in addition to the stresses due to service loading. Tensile residual stresses are detrimental while compressive stresses are beneficial to the rebar performance. In [11], residual stresses on QST rebars were experimentally determined by hole drilling method; compressive residual stresses were found to improve their fatigue strength. In this present study, residual stresses are determined on surface and subsurface of QST rebars by Cut Compliance and X-ray diffraction techniques. Results obtained from both techniques are discussed.

The ribs introduced in the rolling mill as well as the surface imperfections originated from the manufacturing process are zones of stress concentration. The inhomogeneous stress distribution at the ribbed profile and imperfections may initiate a failure and it should be mitigated or if possible avoided.

In this present work, Scanning Electron Microscopy (SEM) analyses were performed on the surface of QST rebars to detect imperfections near the ribs. Similar studies have not been reported in the literature. Moreover, a parametric study was developed to analyse the influence of the rib geometry, including rib inclination, and rebar diameter on the stress concentration factors \( K_t \). \( K_t \) values were determined along the ribs by 3D Finite Element Analyses (FEA)s.

### 3.2 Material characteristics

The chemical composition of QST rebars analysed in this work, with diameters of 16, 26 and 34 mm, is given in Table 3.1. QST rebar is a low alloy carbon steel since it contains a Carbon content \( C < 0.3\% \) and Cu content which can exceed the maximum
Table 3.1 – Chemical composition of the QST rebars with diameters of 16, 26 and 34 mm.

<table>
<thead>
<tr>
<th>Elements</th>
<th>Sample diameter (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>16</td>
</tr>
<tr>
<td>Carbon C</td>
<td>0.186</td>
</tr>
<tr>
<td>Silicon Si</td>
<td>0.22</td>
</tr>
<tr>
<td>Manganese Mn</td>
<td>0.86</td>
</tr>
<tr>
<td>Chromium Cr</td>
<td>0.14</td>
</tr>
<tr>
<td>Nickel Ni</td>
<td>0.13</td>
</tr>
<tr>
<td>Copper Cu</td>
<td>0.40</td>
</tr>
<tr>
<td>Ceq (^2)</td>
<td>0.397</td>
</tr>
</tbody>
</table>

\(^{1}\) It includes only elements with content equal or greater than 0.1%.

\(^{2}\) Ceq: Carbon Equivalent

\[
\text{Ceq} (%) = \text{C} (%) + \frac{\text{Mn} (%) + 0.5 \times \text{Si} (%) + \text{Cr} (%) + \frac{\text{Ni} + \text{Cu} (%)}{4}}
\]

Table 3.2 – Mechanical properties of QST rebars.

<table>
<thead>
<tr>
<th></th>
<th>(\sigma_y)</th>
<th>(\sigma_u/\sigma_y)</th>
<th>(\epsilon_u)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>MPa</td>
<td>MPa</td>
<td>%</td>
</tr>
<tr>
<td>&gt;500</td>
<td>&gt;1.15</td>
<td>&gt;7.5</td>
<td></td>
</tr>
<tr>
<td>&lt;600</td>
<td>&lt;1.25</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Notes:

\(\sigma_y\): yield strength;
\(\sigma_u\): tensile strength;
\(\epsilon_u\): strain at maximum force.

of 0.40% \(^{12}\). Mechanical properties are given in Table 3.2.

3.3 Fabrication process

QST rebars analysed in this work were produced by the Thermex process. Billets with cross section of 170 \(\times\) 130 mm and length of 12.5 m are reheated up in furnaces at about 1200-1300°C. These billets pass through the rollers which squeeze them into the required diameter. After pass by the first rolling the temperature of the bars is between 1140-1180°C. The end rolling temperature is approximately 1050°C. The ribs are introduced on the surface of the bars at the last rolling. The rebars are then slitted in two strands.

Hot rolled steels are then quenched with water in tubes providing intensive cooling of the surface. The rapid quenching leads to the formation of martensite at the
surface layer and a hot core which remains austenitic. The martensite start temperature $M_s$, i.e., the temperature at which the transformation of austenite to martensite begins, can be estimated by Eq. 3.1 [13]:

$$M_s(°) = 539 - 423 \cdot (\%C) - 30.4 \cdot (\%Mn)$$

$$- 17.7 \cdot (\%Ni) - 12.1 \cdot (\%Cr) - 7.5 \cdot (\%Mo)$$

(3.1)

where $M_s$ is approximately 430°C for QST rebars.

When the rebar emerges from the quenching zone, the thermal gradient across the rebar section causes heat to flow from the core toward the surface. This results in a tempered martensitic surface and an equalization of both surface and core temperatures at approximately 670°C [14]. During atmospheric cooling of the rolled bar on the cooling bed, the austenitic core is gradually transformed to a ferrite-pearlite microstructure.

### 3.4 Microstructure

#### 3.4.1 Metallographic analysis

The microstructure of QST rebars was revealed by metallography analysis. Transversal and longitudinal sections were cut with Electrical Discharge Machining (EDM) and polished with abrasive papers and diamond paste [15] up to 0.25 μm. The surfaces were then polished on cloth synthetic suede with alumina up to 0.05 μm for 60 seconds. After polishing, the sections were etched with a fresh solution of Marshall’s reagent [16] for 4 seconds. The sections were completely immersed in this solution where an effervescent etching reaction was observed. The etched surfaces were then rinsed with ethanol, dried with compressed air and analysed under an Optical Microscopy (OM).

Fig. 3.1a shows the transverse etched section of the QST rebar with 16 mm diameter. Fig. 3.1b consists of tempered martensite (TM) within prior austenite grains. Transition zone (TZ) with acicular ferrite and pearlite is shown in Fig. 3.1c and core with quasi-equiaxed ferrite and pearlite (F-P) is given in Fig. 3.1d. The longitudinal section showed similar grain shape to the microstructure of the transversal section in the three TM, TZ and F-P zones.
Figure 3.1 – (a) Etched cross section of the QST rebar; (b) Tempered martensite (TM); (c) Transition zone (TZ) of acicular ferrite (light areas) and pearlite (darker areas); (d) Quasi-equiaxed ferrite (light areas) and pearlite (dark areas) (F-P).

3.4.2 Grain size and area fraction

The Abram’s Three Circle method [17] was used to estimate the ferrite average grain size in the core of QST rebars. Cross sections with diameter of 16, 26 and 34 mm were analysed by OM. The area fraction of pearlite was also estimated according to the procedures given in [18].

The images of the microstructure were taken on the core centre. The average grain size of ferrite and area fraction of pearlite are given in Table 3.3.

The microstructure of the QST rebars became coarser with the increase of the diameter. Similar behaviour was observed by [19] for Tempcore rebars with diameter ranging from 10 to 16 mm.

HR and CW rebars show low (C ≤ 0.3%) or medium Carbon content (0.3% < C < 0.6%) [20, 21]. The microstructure in the entire cross section consists of du-
plex ferrite-pearlite [3, 9]. HR rebars can show coarser microstructure in the core compared to QST rebars with similar or greater diameter. In [9], HR rebars with 12 mm diameter showed an average ferrite grain size of 43 μm which is higher than the values given in Table 3.3. In [22], the average grain size of approximately 25 μm was determined in the centre of HR rebars with diameter of 25 mm.

### 3.4.3 Microhardness test

Hardness may be related to the tensile strength of the steels and it is helpful to determine if the steel has been heat treated in accordance to a specification [10].

Vickers hardness testing was conducted on the QST cross sections with diameters of 16, 26 and 34 mm according to [23]. A hardness map was obtained in the three zones i.e., rim, intermediate and core as given in Fig. 3.1a. A 1 kg-force was applied for 12-15 seconds on the cross sections. The minimum distance of 2.5 t (t is the indenter diagonal) between indentations was respected to ensure that a measurement was not affected by the previous one. This distance was approximately 0.35 mm. The hardness maps are graphically shown in Fig. 3.2.

The minimum and maximum hardness values measured in the rim of the three cross sections were approximately 263 and 285 HV respectively. Similar hardness values can be found in [24] for Tempcore rebars. The hardness decreased continuously from the transition zone to the core where a small variation of the hardness values was measured. As it was expected, the hardness of QST cross sections decreased as the distance from the rim increased.

The core of the cross section with 34 mm showed considerable higher values of hardness compared to both 16 and 26 mm cross sections as shown in Fig. 3.2. This may be explained by the higher amount of pearlite determined in the core of the 34 mm diameter; the hardness increases with increasing pearlite content [25].

### Table 3.3 – Average grain size of ferrite and area fraction of pearlite obtained in the core of QST rebars.

<table>
<thead>
<tr>
<th>Cross section mm</th>
<th>Ferrite ASTM number [17]</th>
<th>95% CI*</th>
<th>Pearlite %</th>
<th>95% CI*</th>
</tr>
</thead>
<tbody>
<tr>
<td>16</td>
<td>11 (7.9 μm)</td>
<td>0.20</td>
<td>11</td>
<td>3.33</td>
</tr>
<tr>
<td>26</td>
<td>10.5 (9.4 μm)</td>
<td>0.19</td>
<td>13</td>
<td>2.04</td>
</tr>
<tr>
<td>34</td>
<td>10 (11.2 μm)</td>
<td>0.30</td>
<td>25</td>
<td>4.62</td>
</tr>
</tbody>
</table>

(*) Confidence interval determined according to the procedures given in [17, 18].

---

36
Figure 3.2 – Hardness map of the QST rebars with diameters of 16, 26 and 34 mm.

CW and HR rebars show an entire cross section of ferrite-pearlite and homogeneous hardness. For HR rebars, the hardness can vary between 160 and 290 HV depending on the steel grade [26, 27].

3.5 Residual stresses

3.5.1 Residual stress on QST rebars

Residual stresses are those which remain in the material with no external applied force. These stresses are in self-equilibrium and consequently the resultant force produced by residual stresses must be zero.

Residual stress pattern on QST rebars result from combination of thermal and microstructural changes in the manufacturing process. The formation of martensite from austenite in the quenching process results in volume expansion at the surface whereas the remainder part is still austenitic. When the remainder austenite transforms to martensite, its volumetric expansion is restricted by the hardened surface layer. This restrain results in tensile residual stress at the surface and compressive residual stress in the interior [10]. At the same time, the interior contraction in the final cooling process is inhibited by the martensitic layer. This restrain generates tensile residual stress in the interior and compressive residual stress at the surface [10]. The tempering process starts just after quenching to minimize these
residual stresses.

Residual stress pattern of rebars are hardly found in the literature. Residual stress acts as a preloading when the rebar is subjected to fatigue loading and it can affect its fatigue strength: compressive residual stresses determined on Tempcore rebars were found to improve their fatigue strength [11] at R=0.

In this work, residual stresses at and near the surface of QST rebars were determined by Cut Compliance (CC) [28] and X-ray diffraction [29, 30] techniques. Since the X-ray diffraction technique is limited to a maximum depth of about 5 μm from the surface of steels, it requires a layer removal at greater depth due to the low levels of penetration of approximately Angstrom wavelength X-ray beams [30].

As-received QST rebars used for residual stress analyses were slightly distorted. This may affect the results obtained from both CC and X-ray diffraction techniques.

### 3.5.2 Cut Compliance technique

CC (also called Crack Compliance) is a destructive technique to determine residual stresses from the material subsurface [28]. A cut is progressively introduced in the specimen using EDM. Strain gauges are glued on the specimen surface and near the cut (see Fig. 3.3) and the strain change is measured at each increment of depth.

![Figure 3.3 – Cut Compliance technique.](image)

The residual stress is numerically estimated from the strain change measured during cutting. Since the cut is assumed as a crack [28], equations of Linear Elastic Fracture Mechanics are used for the residual stress calculation. Stress Intensity factor \( K_{irs} \) due to the residual stress field at the cut tip is given by [31]:

\[
K_{irs}(a) = \frac{E'}{Z(a)} \frac{d\varepsilon}{da}
\]  

(3.2)

where \( \varepsilon \) is the strain measured during the cutting process, \( E' \) is the Young’s Modulus (\( E' = E \) for plane stress and \( E' = E/(1-v^2) \) for plane strain) and \( Z(a) \) is called “influ-
ence function” which depends on the specimen geometry, cut plane and location of the measurement point. Eq. 3.2 is restricted to the Mode 1 case. \( K_{rr}(a) \) is related to the normal residual stresses, \( \sigma_{rr}(y) \), as given in Eq. 3.3.

\[
K_{rr}(a) = \int_0^a f(y, a)\sigma_{rr}(y)\,dy
\]  

(3.3)

Where \( f(y, a) \) is a weight function and \( \sigma_{rr}(y) \) can be obtained by inversion of Eq. 3.3 [31].

**Experimental description**

Residual stresses were determined in a QST rebar with 16 mm diameter. A cut was incrementally introduced at the base of the transversal rib by EDM. Strain changes were measured by surface mounted strain gauges placed about 1.5 mm from the cut rim. Strain measurements were used to determine the residual stresses normal to the cut plane as a function of the cut depth. The measurements were taken near the surface up to 2 mm depth in 40 steps of 0.05 mm.

![Image of a rebar with cuts and strain gauges](image)

**Figure 3.4** – Location and orientation of cuts and strain gauges after measurements.

Position and orientation of cuts and strain gauges after measurements are shown in Fig. 3.4. Cuts were introduced parallel to the uniform rib and also perpendicular to the rebar axis. In this case, the stresses are at the same direction as the fatigue loading on rebars. The elastic constants \( E=205 \) GPa and \( \nu=0.3 \) given in [32] were
used for the residual stress calculations.

**Results and discussion**

The average of the residual stresses evaluated from the strain gauge signals is given graphically in Fig. 3.5. All three profiles showed tensile residual stresses in the region covered by the measurements. The measurements show an imprecision from 5 to 10% up to 1 mm and from 10 to 20% up to 2 mm.

![Residual Stress Graph](image)

**Figure 3.5** – Longitudinal residual stress profile determined on the rebar subsurface.

A peak of stress near the surface is expected in all cases because of the notch effect at the rib base. The stress peak at the surface could not be obtained experimentally since the measurements required the introduction of a minimum cut of 0.05 mm. This peak is expected to be from 10 to 20% higher than the maximum values given at locations 1 and 3 since the rib removal may also release stress. The absence of near surface-stress peak at location 3 may be caused during rib removal by grinding. If the removal was somewhat too deep then the stress peak was eliminated. Moreover, the position of the strain gauge should be at 90° to the cut rim but this angle was not precisely defined.

The combination of these difficulties led to measurement uncertainties. It is likely that the more representative stress peak is at location 2 since the difficulties during the measurements could only reduce the stress peak but not amplify it. Maximum tensile residual stress of 120 MPa can be expected on the surface of QST rebar.
3.5.3 X-ray diffraction technique

Bragg’s law

When X-ray beam irradiates the surface of a crystalline material, with wavelength \( \lambda \) of same order of magnitude as the lattice spacing \( d \) of the material, Bragg diffraction occurs if scattered waves interfere constructively in accordance to the Bragg’s law: the extra path \( 2d \sin(\beta) \) travelled by the deepest wave is equal to integer \( n \) multiples of wavelengths [29] (see Fig. 3.6). When scattered waves satisfy the Bragg’s law, a sharp intensity peak, known as Bragg peak, is produced at those diffracted directions. This intensity peak is observable by the X-ray detector.

\[
n \lambda = 2d \sin(\beta)
\]  

Figure 3.6 – Path difference.

The Bragg’s law is given by Eq. 3.4 [29].

where \( \beta \) is the angle between the incident beam and the diffracting lattice planes. When the elastic properties of the material and the angular position of the Bragg peak \( 2\beta \) are known, then stresses can be calculated [29]. The \( \sin^2 \psi \) method is usually used to determine the residual stresses; \( d \) is plotted versus \( \sin^2 \psi \) and a straight slope \( m \) is fitted by least squares regression [29]. \( \psi \) is the angle between the normal of the specimen and the normal of the diffraction lattice planes.

Experimental description

Residual stresses were determined on the surface and subsurface of a QST rebar with 16 mm diameter. Material was removed by electrolytic polishing for measurements at depths of 0.02 and 0.05 mm. The electro polishing minimizes a possible alteration of the residual stress distribution as a result of material removal [33]. X-ray
measurements were performed between non-uniform ribs at locations MP1, MP2 and MP3 as indicated in Fig. 3.7 and between uniform ribs at locations MP4, MP5 and MP6 as shown in Fig. 3.8.

![Figure 3.7](image1.png)

**Figure 3.7** – Locations of the X-ray measurements between non-uniform ribs.

![Figure 3.8](image2.png)

**Figure 3.8** – Locations of the X-ray measurements between uniform ribs.

CrKa beam was irradiated on the rebar surface over an area between 2 and 3 mm diameter. The X-ray penetration depth was about 5-7 μm. Measurements were performed on the (211) plane with λ of 0.22897 nm. These parameters are recommended for ferritic and martensitic steels [29]. Measurements were taken under four tilt angles ψ of -5°, 5°, -39°, 39° and 2β of 156.4°. Young’s modulus of E=205 GPa and Poisson’s ratio of ν=0.3 were considered for stress calculations. These values are in the range of E and ν values in the (211) atomic plane of steels given in [34]. Residual stresses were determined by the XStress software using the $\sin^2\psi$ method [29].

X-ray elastic constants, such as the Young’s modulus, can produce systematic errors in the measured stresses [29]. Residual stress is proportional to E which can be measured or taken from the literature. However, E values found in the literature don’t always provide the uncertainties around the measured values. Therefore, the uncertainties in E and ν values were set to zero.
3.5.4 Results and discussion

Longitudinal residual stresses determined between uniform and non-uniform transversal ribs are given in Fig. 3.9. Measurement uncertainties varied from 2 to 9 MPa. These uncertainties are related to the peak fitting and the fit of $d$ versus $\sin^2 \psi$ plot.

![Plot of residual stress vs distance from the surface](image)

**Figure 3.9** – Longitudinal residual stresses obtained on the rebar surface and subsurface by X-ray diffraction technique.

Surface compressive residual stresses were obtained in all measurements. They varied from 48 to 147 MPa between non-uniform ribs and from 26 to 61 MPa between uniform ribs. In [2,4], compressive longitudinal residual stress values were also obtained for Tempcore rebars with diameters of 16, 24 and 32 mm. Surface compressive residual stresses of approximately 90 MPa were obtained by extrapolation of near surface measurements performed by the hole drilling method.

Tensile residual stresses varying between 4 to 52 MPa were determined at depth of 0.05 mm from the rebar surface (see Fig. 3.9); only a measurement taken between uniform ribs showed compressive residual stress value of 50 MPa at this depth.

Based on the X-ray diffraction measurements, tensile and compressive residual stresses can be found on the martensitic layer of QST rebars. Since fatigue cracks usually initiate on the martensitic surface layer of rebars and propagates in the presence of tensile stress fields, tensile residual stresses on the surface and near-surface may be detrimental to their fatigue performance.
3.6 Surface imperfections

Imperfections are introduced in the manufacturing process and create a local stress concentration effect (notch effect). They can act as crack initiator and thereby, it is important to quantify them.

In this work, surface imperfections on QST rebars were analysed by XL30-FEG SEM. Five sections were cut, from four as-received QST rebars with 16 mm diameter, by EDM. Size and orientation of these imperfections were determined.

Marks were identified near the transversal ribs as shown in Figs. 3.10a, 3.10b and 3.10c. These marks are introduced in the last rolling mill of the manufacturing process, when the ribs are formed, and they can show semi-circular shape (see Fig. 3.10b).

Cracks were identified near the marks as shown in Figs. 3.10c and 3.10d. Some of these cracks were perpendicular to the rebar axis; their length ranged from few micrometers to approximately 200 µm (see Fig. 3.10d). They may result from the water quenching process which produces rapid cooling and also high residual stresses that can lead to cracks on the material surface [10].

Cracks were also detected near the longitudinal ribs with similar length to the cracks identified at the vicinity of the transversal ribs. These cracks are perpendicular to the longitudinal ribs (see Fig. 3.10e).

![Image](image_url)

**Figure 3.10** – Surface imperfections identified on QST rebars with diameter of 16 mm (a) Marks near the transversal rib; (b) Semi-circular marks near the transversal rib.
**Figure 3.10** – Surface imperfections identified on QST rebars with diameter of 16 mm (c) Marks and cracks near the transversal rib; (d) Cracks near the transversal rib; (e) Cracks perpendicular to the longitudinal rib; (f) Location of the analysed imperfections on the QST rebar surface.
3.7 Stress concentration

3.7.1 Stress concentration on the ribbed profile

A sudden change in the geometrical form of a member section produces local stress distribution. Stress concentration on rebars arises from the ribbed profile (see Fig. 3.11); they can be related to the rib geometry by the stress concentration factor $K_t$. This factor is defined as the ratio between the local stress at the rib and the stress assumed to be uniformly distributed over the total cross section.

![Illustration of the ribbed profile](image)

Figure 3.11 – Illustration of the ribbed profile.

Fatigue performance of rebars can be affected by the stress concentration on the rib geometry. In [22, 27], fatigue life of ribbed bars was considerable lower than smooth bars produced from the same steel. [22] associated the reduction in the fatigue strength of ribbed bars to the stress concentration on the ribs.

2D Finite Element Models FEMs have been used to investigate the influence of the rib geometry on $K_t$ values. [35] concluded that $K_t$ increases as the rib radius $r$ decreases and as the rib width $w$ and flank angle $\alpha$ increases. $K_t$ was found to be independent of the rebar diameter $D$ when $D/h$ ratio ranges from 10 to 20. [36] arrived at similar conclusions as [35]. The highest $K_t$ values were obtained at the root of the ribs.

3.7.2 Numerical analysis

3D Finite Element Model

$K_t$ values were determined for the rib geometry of QST rebars by Finite Element Analyses (FEA)s. Stress analyses were performed using Abaqus/CAE 6.12 software [38]. A parametric study was proposed and a script was developed to generate the 3D rib geometries. The geometrical rib details were provided by the manufacturer. A total of 24 models with different rebar diameters and rib geometries analysed in this paper is given in Table 3.4.

The flank angle $\alpha$ of 47.5° and the shape of the ribs were kept constant in all models. Models without rib radius $r$ were also investigated. Six models represent
Table 3.4 - Geometrical parameters of the analysed models.

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<td>0.15D</td>
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<td>0.6D</td>
<td>D26h15r08</td>
</tr>
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</table>

Notes:
(1) Models which represent the original geometry of the QST rebars. They are referred as original models.
(2) Model with rib inclinations of 90°.
the original geometry of the QST rears i.e., the geometry as provided by the manufacturer. These models are indicated in Table 3.4.

Fig. 3.12 illustrates the typical rib geometry on both sides of the QST rebar analysed in this work. Four ribs were modelled on both sides of the bar and the $K_i$’s were analysed on the centre ribs (see Fig. 3.12). QST rebar shows uniform ribs with inclination $\theta_1$ of $54^\circ$ and non-uniform ribs with inclinations $\theta_2$ of $46^\circ$ and $\theta_3$ of $64^\circ$ as given in Fig. 3.12. Stress analyses were performed for paths 1) along the transition line between the weld toe radius and the rebar cylinder, referred in this paper as TL, and 2) perpendicular to the rib axis as indicated in blue color in Fig. 3.12. The influence of the rib geometry on the bond characteristics between concrete and rebar was not considered in this work. However, the parameter variations of the original models such as rib height $h$ ($0.03D$, $0.15D$) and rib spacing $c$ ($0.4D$, $1.2D$) were chosen according to recommendations given in [37].

![Diagram of rib geometry and paths](image)

**Figure 3.12** – Illustration of the rib geometry and the paths along and perpendicular to the ribs where stress concentrations were investigated.

A free mesh was generated using 10-node quadratic tetrahedral elements ($C_3D_{10}$). The rib radius $r$, which is the critical region on the ribbed profile, was meshed finer. The location away from this region was coarser to save computing time. An uniform tensile stress was applied to the models and the material was considered elastic and isotropic. Material properties $E=205$ GPa and $v=0.3$ were considered as given in [32].

The mesh convergence was verified to obtain an optimum mesh density. Average element sizes $e$ ranging from 0.8 to 0.05 mm were created for $r$ equal to 0.2, 0.4
and 0.8 mm. A resulting mesh is given in Fig. 3.13.

![Typical mesh considered in the models.](image)

**Figure 3.13** – Typical mesh considered in the models.

Fig. 3.14 shows the mesh convergence obtained for \( r = 0.2 \text{ mm} \). \( K_t \) curves obtained along \( r = 0.2 \text{ mm} \) converged for \( e \leq 0.1 \text{ mm} \). The mesh convergence for both \( r = 0.4 \) and 0.8 mm was verified for \( e \leq 0.2 \text{ mm} \).

![Mesh convergence for rib radius \( r = 0.2 \text{ mm} \).](image)

**Figure 3.14** – Mesh convergence for rib radius \( r = 0.2 \text{ mm} \).
3.7.3  \( K_t \) determined on the rib

\( K_t \) values were calculated as the ratio between the largest principal stress and the applied stress. \( K_t \) curves were linearised using a sixth order polynomial function as given in Fig. 3.16. The maximum value obtained in a \( K_t \) curve after linearisation is referred as peak value (PV) in this paper.

Stress concentration profile determined in a perpendicular to the rib axis showed that the zone with higher stress concentrations was obtained at the radius region as shown in Fig. 3.15. PVs determined at the root of the ribs were utmost 20\% higher than PVs determined at TL. \( K_t \) values dropped in direction to the centre of the rib where no stress concentration was observed.

![Stress concentration profile](image)

**Figure 3.15** – Stress concentration profile obtained in a perpendicular to the rib axis of D26ho8ro2 model.

As show in Fig. 3.15, the rib itself does not show high stresses. The radius region was the critical region since it showed the maximum stress concentrations. This is in agreement with the results of [36].

**Influence of the rib inclination**

Fig. 3.16 shows typical \( K_t \) curves obtained along TL. The smallest stresses along PT were obtained at the rib ends. The maximum \( K_t \) value at these locations was 1.2. PVs were similar between the different rib inclinations \( \theta \). However, the peak positions (PP)s tended to move to the centre of TL with the increase of \( \theta \); for \( \theta \) of 90\° for
example, PP was at the centre of TL as shown in Fig. 3.16.

\[ K_t \]

![Graph](image)

**Figure 3.16** - \( K_t \) curves obtained for different rib inclinations \( \theta \) of D16h08r04 and D16h08r04θ90 models.

PVs and PPs determined along TL for different \( \theta \) are given in Table 3.5.

Zones along TL with \( K_t \) equal or higher than 95\% of PVs were analysed. These zones are referred as peak zones (PZs) in this paper. PZs of the original models were anti-symmetrically located along TL and they showed similar sizes as given in Fig. 3.17. Fig. 3.18 shows PZ sizes obtained for different \( \theta \). PZ sizes increased along TL as \( \theta \) increased; \( \theta \) of 64°, for example, showed higher PZ sizes compared to the ribs with inclination of 46° and 54°. However, the increase of \( \theta \) from 64° to 90° had a negligible influence on the PZ sizes.

**Influence of the radius**

The increase of \( r \) resulted in the reduction of the PVs of the original models (see Table 3.5). PVs decreased in 14 to 21\% with the increase of \( r \) from 0.2 to 0.4 mm and dropped by 15\% with the change of \( r \) from 0.4 to 0.8 mm. The maximum PV obtained along TL for the original models was 2.50. In the models with no radius, PVs was utmost 20\% higher than PVs of the models with \( r=0.2 \) mm.

Fig. 3.19 shows the PPs and PZs obtained along TL with the change of \( r \). Small variation on the PPs of the original models was obtained with the increase of \( r \). The increase of \( r \) led to an increase of the PZ sizes: the change of \( r \) from 0.2 to 0.4 mm,
for example, resulted to an increase up to 11% on the PZ sizes.

**Influence of the height**

The reduction of \( h \) resulted in the decrease of PVs as shown in Table 3.5. The change of \( h \) from 0.075\( D \) to 0.03\( D \) reduced PVs from 14 to 21%. The maximum PV determined for the models with \( h=0.3D \) was 2.09. Fig. 3.20 shows the PPs and PZs obtained for models with different \( h \). Small variation on the PPs was observed; PZ sizes increased up to 15% with \( h \) reduction from 0.075\( D \) to 0.03\( D \). The increase of \( r \) from 0.2 to 0.4 mm had similar effect on PVs as the change of \( h \) from 0.075\( D \) to 0.03\( D \).

The maximum PV of 2.79 was determined between all analysed models (see

---

**Table 3.5** – Peak values (PV) and Peak positions (PP) determined along the transition line (TL).

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**Notes:**
- D1o8084020 model: all ribs with the same inclination of 90°;
- PV = 1.85 and PP = 0.5.
Figure 3.17 – Example of the peak positions (PP)s and peak zones (PZ)s along the rib of an original model.

Figure 3.18 – Influence of the rib inclination $\theta$ on $K_t$.

Table 3.5). This value resulted from a combination of highest $h=0.15D$ and smallest $r=0.2$. Models with $h=0.15D$ combined with $r=0.4$ or 0.8 mm affected both PVs and
PPs of the non-uniform ribs. As the flank angle $\alpha$ was kept constant in all models, the increase of $h$ was followed by the increase of the rib width $w$. Beside, the increase of $r$ and the different rib inclination $\theta$ reduced significantly the distance between the non-uniform ribs. As a consequence, it created a region of stress concentrations moving the PPs of the ribs to this zone. The reduction of the distance between non-uniform ribs increased PVs of these ribs. Therefore, these rib geometries should be avoided.

**Influence of the diameter**

The increase of $D$ resulted in the increase of PVs as shown in Table 3.5. PVs of the original the models increased up to 14% as $D$ changed from 10 to 16 mm and to utmost 18% for $D$ increasing from 16 to 26 mm.

Fig. 3.21 shows PPs and PZs obtained for the original models with 10, 16 and 26 mm diameters. Small variation on the PPs and PZ sizes was obtained with the change of $D$ from 10 to 16 mm and from 16 to 26 mm.

**Influence of the rib spacing**

Rib spacing $c$ change had small influence on the PVs and PZ sizes along TL as shown in Fig. 3.22. The $c$ parameter had smaller influence on the $K_I$ values and PZ sizes.
compared to other analysed parameters.

However, the reduction of $c$ from 0.6$D$ to 0.4$D$ affected the PVS and PPs of the non-uniform ribs (see Table 3.5). Similar case was obtained for models with $h=0.15D$ combined with $r=0.4$ or 0.8 mm.
3.8 Conclusions

In this work, characterisation studies of QST rebars included 1) Identification of the microstructure and hardness measurements in the cross section of rebars with different diameters. The average grain size of the core microstructure was quantified and compared to the grain size of QST and HR rebars given in the literature. 1) Residual stresses were experimentally determined on surface and subsurface of QST rebars with 16 mm diameter by Cut Compliance and X-ray diffraction techniques. Results obtained from both techniques were discussed and compared to residual stresses of QST rebars found in the literature. 2) SEM analyses on the rebar surface allowed to identify and quantify imperfections originated in the manufacturing process. 3) A parametric study was developed to analyse the influence of the rib geometry and rebar diameter on the stress concentration factors $K_I$ along the rib. The findings of this study suggest that:

- The average grain size in the core of QST rebars increase with the increase of diameter. However, the microstructure of QST rebars is finer than the microstructure of hot rolled rebars with higher diameter.

- Random tensile and compressive residual stresses are expected on the surface and subsurface of QST rebars. These stresses are no higher than 20% of their yield strength.
• Surface imperfections originated in the manufacturing process are near the transversal and longitudinal ribs. As imperfections are zones of stress concentration and may affect the performance of rebars in service, they should be avoided.

• The highest stress concentrations arising from the ribbed profile are located at radius region. The main parameters of the rib geometry affecting the maximum $K_i$’s are the radius and height; maximum $K_i$ values at the radius region can be significantly reduced with the increase of the radius and reduction of the height.

• The rib inclination is the main parameter affecting the position of the maximum $K_i$’s as well as the critical zone sizes along the rib. The critical zones correspond to the zones where $K_i$’s are at least 95% of the maximum values. These critical zones are located anti-symmetrically along the rib. As near the rib inclination is to a perpendicular of the rebar axis, the maximum $K_i$’s tends to move to the rib centre and the critical zone sizes increase.

• Maximum $K_i$’s along the ribs increase with the rebar diameter increasing and its influence in the maximum $K_i$ is comparable to the changes in the rib radius and height. However, the effect of the diameter on the critical zone size is less significant than those parameters.

• The reduction of the rib spacing should be avoided for non-uniform ribs since it increases the maximum $K_i$ values.

3.9 Acknowledgements

The authors are grateful to Danièle Laub from the Interdisciplinary Centre For Electron Microscopy (CIME) at EPFL for her help with the sample preparation for the microscopic analyses. We are also grateful to Cyril Dénéraz from the Mechanical Metallurgy Laboratory (LMM) for the hardness measurements.
Bibliography


Microstructural influence on the scatter in the fatigue life of steel reinforcement bars
— Marina Rocha, Eugen Brühwiler, Alain Nussbaumer

Abstract: Fatigue test results with steel reinforcement bars (rebars) is a stochastic process. Scatter can be influenced by the sensitivity of the short crack growth to the microstructural features, especially near the fatigue limit. This work investigates the scatter inherent to the microscopic conditions near the fatigue limit of ferrite-pearlite and martensite microstructures found in the outer layer of rebars. An adapted Navarro-De Los Rios model within a Monte-Carlo framework is used to simulate the short crack growth in material grains. Grain size variation, grain orientation factor and multiple phases were considered in the model. The results are compared with the scatter found in fatigue tests on hot-rolled-cold worked as well as quenched and tempered rebars. It is shown that microstructural effects explains part of the observed scatter in the fatigue tests.

Keywords: Short fatigue crack growth; Ferrite-pearlite; Tempered martensite; Scatter above the fatigue limit.
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### 4.1 Introduction

Fatigue life assessment of engineering structures depends on the knowledge of material properties. Fatigue limit is a macroscopic property of the material that is sensitive to microscopic conditions. Since metal fatigue is a random process, fatigue data from testing always exhibit scatter for specimens of the same geometry, under same loading condition, especially near the fatigue limit. Besides, for metallic materials, the crack initiation period often dominates the fatigue life just above this limit. Since fatigue crack initiation is mainly a surface phenomenon, specimen surface roughness, protrusions and microscopic aspects such as grain size, grain orientation, dislocation density can influence the scatter in fatigue lives and fatigue limit stress range. The large scatter observed in fatigue tests lead to high values of confidence intervals (CI)s to get the design S-N curves. Further scatter result from the damage accumulation models used in fatigue life verifications. In most design codes, including those for steel reinforcement bars (rebars), usually considers 95% CIs.

Furthermore, one single design S-N curve regroup rebars produced from hot rolling, cold working and quenching and self-tempering with different surface microstructures, rib patterns and surface roughness, all of which increase the scatter near the fatigue limit and lead to very conservative design values. In order to understand the scatter inherent to microscopic aspects in the fatigue behaviour of rebars, this work proposes a short crack growth model to simulate the initiation period
in ferrite-pearlite and tempered martensite grains found on the surface of different types of rebars.

Fatigue crack initiation period in metallic material consists of cyclic slip, crack nucleation and short crack growth [1]. These cracks can behave significantly different from the (long) crack propagation as described by Paris’ law [2,3]. Short cracks can grow faster than corresponding long cracks at the same value of stress intensity factor range, $\Delta K$. They can also grow at significant rates for $\Delta K$ value smaller than the threshold for long cracks [4].

Surface microscopic investigations in steel specimens has shown that short cracks form on slip bands and propagates along them [3]. The short crack propagation stops when the applied stress is below the fatigue limit; when the stress is just above this limit, the crack accelerates and decelerates due to the interactions with microstructural barriers until it reaches a long crack regime with apparent continuous propagation rate. The crack growth rate decreases as approaching the grain or phase boundary; the crack then accelerates when a slip band is initiated in the adjacent grain [2,5].

Navarro and De Los Ríos (N-R) [6,7] proposed a short crack growth model where a crack initiates from slip bands. This model is particularly appropriate to study fatigue limit problems which involve microstructurally short cracks. N-R model considers the interaction between crack and microstructural barriers. It assumes that dislocations are constrained to remain on their original plane and pile up when blocked by grain boundaries. They propagate along slip bands extending through successive grains.

N-R model can be considered as a further development of research in [3,8,9]. Bilby et al. [8] were the first to obtain a bounded solution for the dislocation distribution function representing the crack and the plastic zone (slip band ahead of the crack tip). [3,9] showed that the crack tip plastic zone interacts with microstructural barriers, such as grain boundaries, when the crack is of the order of microstructural features. [9] developed the unbounded solution for the dislocation distribution function.

In its original form, N-R model considers infinitesimal dislocations distributed within two zones: the crack itself $a$ (half crack length) and the plastic zone $c$ as shown in Fig. 4.1. In this two-zone model, an infinite stress level is sustained by the grain boundary. In [10], the model was extended by considering an additional small zone of length $r_0 << D$, ($D$ is the, uniform, grain diameter) representing the interface between neighbouring grains or phases (see Fig. 4.1). This model elimi-
nates the singularity of the stress field associated to the distribution of dislocations. In physical terms, the three-zone system was argued to be more realistic since the plastic zone is blocked by the grain boundary i.e., its two boundaries, and it remains blocked until the stress in the third zone i.e., the grain boundary, attains a critical level for dislocations to cross this zone.

![Diagram of Two-zone and Three-zone models](image)

**Figure 4.1** – Two-zone [6], [7] and three-zone [10] models.

N-R model has been extensively used (or extended) to predict the fatigue lifetime of metals. In [11], N-R model was applied to predict the short fatigue crack growth behaviour in mild steel. In [12], N-R model was applied for fatigue life prediction of commercially pure aluminium. The model prediction was in good agreement with experimental results. In [13], the effect of textures was investigated in the short fatigue crack growth in Al-Li alloy. An equation for the grain orientation factor was proposed depending on the load axis, slip plane normal and slip direction. N-R model was extended for biaxial fatigue loading case in low and medium Carbon steels; the crack initiation orientation was close to the experimental results [14]. The influence of the grain size variation was introduced in the N-R model by [15] where Voronoi cells were used to represent the grain structure. In [16], a micromechanical model for short crack growth based on successive blocking of monotonic plastic zone and cyclic plastic zone of a crack at grain boundaries was proposed. This model was based on the N-R approach [10]. These models successfully reproduced the short crack growth pattern where the crack decelerates at a grain boundaries and it accelerates when crossing a grain.
This irregular behaviour of short cracks due to the interactions with the microstructure can affect the scatter found near the fatigue limit. Factors such as grain size variation [3], grain orientation [13] and different phases [2] are more prominent at low stress levels and consequently influence the fatigue crack initiation period.

In this study, the influence of the microstructure on the scatter observed in experimental data as obtained above the fatigue limit is investigated using an adapted N-R model within a Monte-Carlo framework. The short crack growth is simulated in ferrite-pearlite (F-P) and tempered martensite (TM) grains found on the outer layer of rebars. The grain structure is represented by Voronoi tessellation. In the F-P model, the two phases, pearlite and ferrite, were modelled separately with the area fraction of each phase being obtained from the literature. On the other hand, the parent austenite grain is considered for short crack growth modelling in TM. The parent grain is experimentally determined using Electron Back Scattered Diffraction (EBSD) and ArpGe software. The scatter obtained in the crack initiation phase is compared to experimental data from the literature.

4.2 Reconstruction of parent austenite grains

The laths of (daughter) martensite are organized within the parent austenite grains. The austenite grain size represents an essential characteristic of martensite steels; fine austenite grain results in the formation of fine martensite and consequently improvement in the mechanical properties of martensite steels [17]. In this work, slip band and short crack growth is simulated in the austenite grains of TM steel.

With the martensite transformation, also called displacive transformation, a vestige of the austenite grain boundary remains in the microstructure [18] and it can be revealed by etching [19]. However, it can be time consuming to find a successful etching to correctly identify the austenite grain boundaries. Instead, in [20], parent grains were successfully reconstructed by post-processing of EBSD data on the daughter grains. This technique was applied in this work to reconstruct the parent austenite grains in the cross section of quenched and self-tempered (QST) rebar from EBSD data obtained on martensite laths. The ArpGe software [21] was used for reconstruction of the parent grains.

4.2.1 Experimental analysis

EBSD analyses were performed on a polished cross section of QST rebar with 16 mm diameter. The cross section was prepared by mechanical grinding and polishing
Figure 4.2 – Reconstruction of the austenite grains in the QST rebar: (a) Martensite grains obtained from EBSD analyses; (b) Reconstructed austenite grains determined from ARPG software.
up to 1 μm followed by polishing with a Vibromet table containing non-crystalline colloidal silica for approximately 3.5 hours. EBSD analyses were then conducted using XL30-FEG Scanning Electron Microscopy (SEM) at 20 KV. The cross section was tilted at 70° in the SEM. EBSD map of the martensite grains was obtained from the surface edge to 200 μm towards the centre. A map size of 200 × 200 μm was considered with a measurement step size of 0.2 μm.

EBSD map of the martensite laths is given in Fig. 4.2a. The orientation of the martensite grains is coded by colors representing the Euler angles; they are a set of three angles which describe the crystallographic orientation of grains relative to a reference (sample) coordinate system.

The austenite grains reconstructed by the ARPEG software are shown in Fig. 4.2b. The reconstruction was performed in three steps: identification of the martensite grains in the EBSD mapping; nucleation of austenite grains based on a selected orientation relationship (OR); growth of the austenite grains up to a defined tolerance angle [20]. The OR between austenite and martensite was the Greninger-Troiano (GT) relationship [18]. The white areas in the image (see Fig. 4.2b) correspond to the areas that could not be reconstructed by ARPEG. This may result from no indexed pixels in the EBSD map, martensite with ORs that are far from the GT-OR, martensite laths that could belong to two parent austenite grains and for which it was impossible to decide [20]. The color of each austenite grain represents the color of one martensite lath into this grain.

4.3 Short crack growth model

The short fatigue crack growth model is adapted from the three-zone micromechanical N-R model [10]. The interaction between crack and grain-phase boundary is characterized. It is assumed that when slip is initiated in a grain, the whole grain undergoes slip being blocked by the grain boundary. Slip propagates to the adjacent grain when the stress ahead of the plastic zone reaches a required value. This stress depends on the position of the crack tip to the grain boundary.

4.3.1 Grain structure

In N-R model, all grains are assumed to have equal sizes. In this work, the grain structure with different sizes was represented by Voronoi tessellation. Voronoi has been extensively used to reproduce the grain geometry of polycrystalline materials in short fatigue crack initiation models [15, 22].

70
Figure 4.3 – Example of a simulation result obtained in F-P with a crack length $2a \approx 8D_m$ at $N = 2 \times 10^6$ cycles: (a) Illustration of a surface short crack growth in ferrite (dark grey) and pearlite (light grey) grains represented by Voronoi cells; (b) Short crack growth rate as a function of the crack length.
Fig. 4.3a illustrates a short crack in F-P steel where the two phases, ferrite and pearlite, are modelled separately by Voronoi cells. The crack grows along a preferential slip direction randomly distributed in the model. In TM steel, Voronoi cells represent the parent austenite grains.

Voronoi tessellation is a cell structure generated from a random set of points as shown in Fig. 4.3a. Voronoi tessellation is a good approximation to represent a polycrystalline structure assuming that, in the crystallization process of a metal, all grains start to grow from random points and grow uniformly in all directions. The grains will then collide and a grain boundary will be created. In geometric aspects, the resulting grain structure would be a Voronoi.

In the Voronoi tessellation, the set of points is specified beforehand and each point will be enclosed by an area. Each Voronoi area (or cell) represents one grain. Voronoi tessellation was generated using MATLAB.

**Crack initiation in the first grain**

Short fatigue cracks in steel specimen under uniaxial loading initiate along slip planes closer to the plane of maximum shear stress [24]. The angle between the slip plane and the applied load axis is close to $45^\circ$ (mode II loading). Moreover, crack initiates in relatively coarse grains in steel [2]. Thus, it was assumed in the model that a crack initiates in a large grain along the slip plane closer to the plane of maximum shear stress. The crack always starts from a Voronoi point. To decide in which grain the crack will start, a compromise between slip size and direction was considered. A new length $l_{cr}$ was calculated for each grain as given in Eq. 4.1:

$$l_{cr} = l \cos 2\theta$$

(4.1)

where $l$ is the length of all slip planes passing through the Voronoi point and $\theta$ is the angle between the slip plane and the plane of maximum shear stress. The crack initiates in the grain with maximum $l_{cr}$.

**Short crack growth**

The schematic diagram given in Fig. 4.4 represents a half crack length $a$ and its plastic zone $c$ in a polycrystalline material. The crack grows in mode II loading and the applied shear stress component is assumed as $\Delta\tau = \Delta\sigma/2$, where $\Delta\sigma$ is the applied uniaxial cyclic loading. In this model, a single crack is assumed to propagate until it reaches a given length of $\sigma D_{mean}$ ($D_{mean}$ is the mean grain diameter) [2, 25]
and the simulation is stopped. $D_{\text{mean}} = 20 \mu m$ as obtained from Fig. 4.2b using the Abram's Three Circle method [23].

The crack growth is calculated for each crack tip separately. The crack grows at a rate of:

$$\frac{da}{dN} = f \phi$$

(4.2)

where $N$ is the number of cycles, $f$ is the fraction of dislocations ahead of the crack tip that contributes to the crack growth process and $\phi$ is the crack tip plastic displacement.

The factor $f$ varies between 0 and 1 and it depends on the applied stress: the smallest $f$ values are obtained when the applied stress level approaches the fatigue limit [24]. It represents the degree of irreversibility of slip per cycle. In this work, $f$ was assumed constant and equal to $5.64 \times 10^{-4}$ since the scatter is analysed at constant stress amplitude and near the fatigue limit. This value was obtained from fatigue tests with low carbon steels subjected to uniaxial loading [24].

The plastic displacement $\phi$ can be written as [26]:

$$\phi = \frac{2(1 - v)^2 \sqrt{1 - n^2} \Delta \tau}{\mu n}$$

(4.3)

where $v$ is the Poisson's ratio and $\mu$ is the shear modulus. The dimensionless parameter $n = a/c$ defines the position of the crack tip relative to the grain boundary (see Fig. 4.4).
The crack growth rate as a function of the crack length obtained for F-P is given in Fig. 4.3b. The growth rate decreases as the crack approaches the grain boundary and then increases as the plastic zone propagates into the next grain. The dotted lines in Fig. 4.3b represent the plastic zone size at every propagation step in each grain.

The slip propagates from grain \( i \) to the neighbouring grain \( i + 1 \) when \( n \) reaches a critical value \( n = n_c \). This critical value \( n_c \) is given by [10]:

\[
n_c^i = \cos \left( \frac{\pi}{2} \left( \frac{\Delta \tau - \Delta \tau_{Li}}{2 \tau_{fr}} \right) \right)
\]  

(4.4)

where \( \tau_{fr} \) is the "friction stress" which represents the resistance of the grain to the dislocation motion in the plastic zone. \( \Delta \tau_{Li} \) is the minimum stress required for slip propagation determined as [15]:

\[
\Delta \tau_{Li} = \Delta \tau_{FL} \frac{m_i}{m_1} \sqrt{\frac{d_i}{2c_i}}
\]  

(4.5)

where \( \Delta \tau_{FL} \) is the fatigue limit, \( d_i \) is the mean of the crack length in each grain and \( m_i/m_1 \) is the grain orientation ratio. \( \Delta \tau_{FL} \) represents a macroscopic property of the material and \( \Delta \tau_{Li} \) represents a condition for local plastic deformation at the microscopic level. The \( m_i/m_1 \) ratio will be discussed in Section 4.3.2. The crack propagation stops if \( \Delta \tau \) is smaller than \( \Delta \tau_{Li} \).

When a slip band starts in the next grain, the plastic zone spans this entire grain. Thus, \( n_c^i \) decreases to \( n_c^{i+1} \) expressed as:

\[
n_c^{i+1} = \frac{c}{c_i+1} \cdot n_c^i
\]  

(4.6)

The number of cycles to propagate the crack in a grain can be obtained by integration of Eq. 5.3 which gives:

\[
\Delta N_i = \frac{\mu}{f(1-v)\Delta \tau} (\sin^{-1}n_c^i - \sin^{-1}n_c^{i+1})
\]  

(4.7)

The number of cycles \( N_{d1} \) and \( N_{d2} \) spent to propagate each crack tip separately are determined from the sum of \( \Delta N_i \). The total number of cycles \( N_{Total} \), when the crack length is equal to \( 10D_{mean} \), is given by:

\[
N_{Total} = \max(N_{d1}, N_{d2})
\]  

(4.8)
4.3.2 Grain orientation

An equation for the grain orientation ratio \( m_i/m_1 \), used in N-R model, was proposed by [27] and it is given as:

\[
\frac{m_i}{m_1} = 1 + 2.07 \left[ \frac{2}{\pi} \tan^{-1}\left(0.522(i - 1)2\right) \right]^{1.86}
\] (4.9)

where \( m_1 \) is the orientation factor of the first grain and \( m_i \) is the orientation factor of successive grains. Eq. 4.9 was developed based on fatigue test results of mild steel and then applied for randomly oriented fcc materials. \( m_i/m_1 \) increases progressively as the crack grows over several grains (see Fig. 4.5b) and \( m_{1 \to \infty} \) tends to the average Taylor factor \( \tilde{M}_T = 3.07 \).

In reality, the short crack growth depends on the individual grain orientations. Since the objective of the present work is to analyse the scatter due to microstructural features including the grain orientation, a new equation \( m_i/m_1 \) was proposed where this factor isn’t forced to increase continuously with the crack length; it can decrease or increase as the crack propagates through successive grains which are more or less favourably oriented for slip. This allows to consider how the statistical variations of the grain orientation can affect the crack growth.

\( m_i/m_1 \) orientation ratio

The crack initiation in the first grain is governed by the stress required to sustain plastic deformation along the slip direction with the most favourable orientation, which results in the smallest orientation factor. This factor measures the deformation compatibility between the current crack and the slip direction in the adjacent grain. Since the need to maintain the compatibility between adjacent grains in this stage is small, the orientation factor is close to the Sachs factor for single crystal. This factor \( \tilde{M}_S \) is equal to 2.24 (Sachs factor) for fcc and body centered cubic (bcc) materials [28].

With further crack propagation over several grains with different orientations (see Fig. 4.5a), the deformation compatibility between all these grains moves the orientation factor value from Sachs to Taylor type. When the crack becomes sufficiently large i.e., insensitive to the material microstructure, the orientation factor reaches a maximum value similar to the Taylor factor, \( m_{1 \to \infty} = \tilde{M}_T = 3.07 \) for randomly oriented fcc and bcc crystals [28].

Theoretically, the grain orientation ratio \( m_i/m_1 \), for \( m_1 = 2.24 \) and \( m_{1 \to \infty} = 3.07 \),
Figure 4.5 – Illustration of a subsurface crack growth and variation of the grain orientation ratio $m_i/m_1$: (a) A crack length $2a$ reaches 7 grains on the surface, the subsurface crack tip extends over $N = 14$ grains; (b) The graph represents the evolution of $m_i/m_1$ scatter as the crack grows using Eq. 4.10; To illustrate this scatter, it was computed 1000 $m_i/m_1$ values for each step. The greyscale represents the amount of $m_i/m_1$ with same values in a region. In the first step, $m_i/m_1 = 1$ and there is no dispersion. In the second step, a higher dispersion is obtained compared to the 10th step when the dispersion decreases. The red lines represent the evolution of $m_i/m_1$ obtained by Eq. 4.9.
varies between 1 and 1.4. However, as given in [29, 30], \( m_{i-\infty} / m_i \) can tend to a value of approximately 3 for mild steels. [29] suggests that this difference is attributed to crack closure effects; as the crack grows, it progressively increases to a steady peak value for large crack. Other factors such as microcrack initiation at several points and microcrack coalescence may also contribute to the value of 3.

Eq. 4.10 gives \( m_i / m_i \) which simulates the variance for steels with randomly oriented crystals:

\[
\frac{m_i}{m_i} = 1 + 2(\Omega_i - 2.3) + 2.07\left[ \frac{2}{\pi} \tan^{-1}\left(0.522(i - 1)2\right) \right]^{1.86} \tag{4.10}
\]

where \( \Omega_i \), the average value of the grain orientations, is calculated as:

\[
\Omega_i = \frac{1}{N} \sum_{j=1}^{N} \omega_j \tag{4.11}
\]

\( N \) represents the number of grains which contains the crack tip as shown in Fig. 4.5a.

\[
\omega_j = \min_{k=1\text{ to } 12} \left( \frac{1}{(\mathbf{L} \cdot \mathbf{n}_k)(\mathbf{L} \cdot \mathbf{s}_k)} \right) \tag{4.12}
\]

where \( \mathbf{L}, \mathbf{n}_k \) and \( \mathbf{s}_k \) are unit vectors along the loading axis, slip plane normal and slip direction respectively. The \( k \) values represent the main slip system in bcc crystals: 6 slip planes and 2 slip directions in each plane.

Fig. 4.5b shows the \( m_i / m_i \) behaviour obtained for 1000 simulation runs at each step of the short crack growth. When the crack propagates in the first grain, the grain orientation ratio is 1. As the crack grows in few grains, \( m_i / m_i \) variation increases due to the higher dispersion in grain orientations. As more grains are incorporated to the crack growth, this variation decreases and \( m_i / m_i \) tends to the value of 3 when the crack length \( 2a \approx 10D_m \).

### 4.3.3 Material properties

The parameters used for ferrite, pearlite and martensite in the model are given in Table 4.1.

#### Ferrite and pearlite properties

Fatigue limit \( \Delta \sigma_{FL} \) of ferrite steels, i.e., with very low amount of pearlite, is approximately 0.6\( \sigma_u \) [31], where \( \sigma_u \) is the tensile strength. This results in \( \Delta \sigma_{FL}=260 \)
Table 4.1 – Material properties used in the model.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Ferrite</th>
<th>Pearlite</th>
<th>Martensite</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\tau_f$ (MPa)</td>
<td>70</td>
<td>134</td>
<td>142</td>
</tr>
<tr>
<td>$\Delta\sigma_{FL}$ (MPa)</td>
<td>260</td>
<td>385</td>
<td>368</td>
</tr>
<tr>
<td>$\mu$ (GPa) [40]</td>
<td>82</td>
<td>82</td>
<td>79</td>
</tr>
<tr>
<td>$\nu$ [40]</td>
<td>0.28</td>
<td>0.28</td>
<td>0.29</td>
</tr>
</tbody>
</table>

MPa [32]. $\tau_f$, for ferrite is 70 MPa as given in [33].

Fatigue limit $\Delta\sigma_{FL}$ of fully pearlite steels is approximately $0.4\sigma_u$ [31]. The value of 385 MPa was obtained [34] as shown in Table 4.1. The $\tau_f$ value of 134 MPa is given in [35]. The area fraction of pearlite for hot rolled (HR) rebars was found to vary from approximately 15 to 55% [36, 37]. This variation was considered in the model.

Since in fatigue tests with F-P steels, cracks usually start in the softer ferrite grain [24, 38, 39], the crack was assumed always to start in a ferrite grain in the model.

Tempered martensite properties

Fatigue limit $\Delta\sigma_{FL}$ of TM steels was found to be approximately $0.6\sigma_u$ [31]. Since $\sigma_u=613$ MPa for the TM steel studied in this work, $\Delta\sigma_{FL}$ was then determined as approximately 368 MPa (see Table 4.1). The $\tau_f$ values given in [41, 42] for TM steels varies from approximately 0.25 to 0.30$\sigma_y$ with $\sigma_y$ being the yield strength. $\tau_f = 142$ MPa was calculated from $0.275\sigma_y$ with $\sigma_y = 518$ MPa for the studied TM steel.

4.4 Results and Discussion

Fig. 4.6 shows the scatter obtained in the F-P model for different area fractions of pearlite i.e., varying approximately between 15 and 55%, as found in HR rebars [36], [37]. The scatter in F-P model is affected by the presence of two phases and different area fractions of ferrite and pearlite. As the crack grows into pearlite from softer ferrite phase, the crack growth rate decreases at phase interface. Since pearlite is a harder microstructure, the boundary between two phases tended to impede the crack propagation. Similar crack behaviour was experimentally observed in fatigue tests with F-P steels [38]. As the area fraction of ferrite varies considerably between HR rebars, the duration of fatigue crack initiation was affected with this area variation and consequently, the scatter at the analysed stress level. Smaller area fraction
Figure 4.6 – Scatter obtained for different area fractions % of pearlite in the F-P model.

of pearlite, for example, tended to decrease scatter since the crack grows predominantly in one phase i.e., in ferrite grains.

Figs. 4.7a and 4.7b show the scatter determined from simulations of the short crack growth in F-P (with 53% of pearlite) and TM steels. The model results were compared to experimental data including rebars with diameter $d \leq 16$ mm and tested at stress-ratio between 0 and 0.2 as given in [43–47]. Model and experimental results given in Figs. 4.7a and 4.7b include only failure data points. The model results include the influence of microstructural features on the scatter in the fatigue crack initiation phase. This phase includes short crack propagation to a maximum length of 0.2 mm. The long crack propagation phase will be treated in another paper.

The model data were generated from 1000 simulations run at 24 MPa above the fatigue limit of both F-P and TM steels. The scatter at this stress level was compared to the scatter found in HR-CW and QST rebars with similar local conditions. The value of 24 MPa was considered in the simulations since 1) The smallest $f$ value experimentally given in [24] was determined for a stress range at 24 MPa above the fatigue limit of the material; 2) There were no or few experimental data for HR-CW and, especially for QST rebars, at lower stress ranges and with the same test conditions as considered in this work.

The experimental data in Fig. 4.7a include HR and CW rebars of low and medium Carbon content with different area fractions of pearlite. Information on the crystallographic texture of HR, CW and QST rebars isn’t provided in the literature. Crystal-
Figure 4.7 – Distribution of failures points obtained in the model and experimental data: (a) For F-P model with an area fraction of 53% of pearlite and HR-CW rebars [43, 44]; (b) For TM model and QST rebars [45–47].
lographic texture is defined as the grain (crystal) orientation in a specimen. In [48], nearly random texture was found on the surface of hot rolled low-alloyed steels with bcc structure; during hot rolling, the steel undergoes several passes where the texture is randomized by dynamic recrystallization and phase transformation from austenite to ferrite. Randomly oriented grains were then assumed for F-P and TM in the model as a first approximation.

Fig. 4.7b shows the scatter of failure data obtained from the model and QST rebar test results. With the model, higher scatter was obtained in F-P compared to TM which is in agreement to what has been observed in fatigue tests with HR, CW and QST rebars. In [49], for example, QST rebars showed smaller scatter compared to HR and CW rebars. In [45], small scatter was observed in fatigue tests with QST rebars near the fatigue limit. The fatigue behaviour of these rebars was considered essentially as a function of the surface layer of martensite.

Moreover, as shown in Figs. 4.7a and 4.7b, the scatter determined in both F-P and TM is smaller than the scatter observed in experimental data; it represents about 50% of the scatter from fatigue tests. This difference between model and experimental results may be influenced by stress concentrators on the rebar surface, such as roughness and the presence of the ribs, giving rise to local plastic deformation and affecting fatigue crack propagation. [50], for example, suggests that scatter near the fatigue limit of QST rebars may result from variations in stress concentrations arising from ribs as well as surface microstructural features.

### 4.5 Conclusion

N-R model including grain structure with different sizes and grain orientation variations was used to simulate the propagation of a single fatigue short crack in ferrite-pearlite and tempered martensite steels within a Monte-Carlo framework. The scatter determined above but near the fatigue limit was compared to the scatter found on experimental data of rebars. Grain size variation using Voronoi tessellation, randomly oriented grains and different phases were considered in the analyses. This work suggests that:

Martensitic microstructure contributes to the scatter reduction in rebars at stress range levels above the fatigue limit compared to ferrite-pearlite. The grain and phase boundaries as well as different area fractions of ferrite-pearlite affects the short crack growth rate and thus increase the scatter in the fatigue crack initiation period.

The microstructure on the rebar surface is responsible for about half of the scat-
ter found in fatigue tests with rebars used as reference. Since fatigue crack initiation is mainly a surface phenomena, it is likely that surface roughness and ribs may also contribute to the dispersion in the tests.

For further studies, it is recommended to consider:

- More experimental data, especially on QST rebars, is needed near the fatigue limit. Few data were found in the literature with similar test conditions as considered in this work.

- The crystallographic orientation of hot rolled, cold worked and quenched and self-tempered rebars, considered in this work, should be confirmed. Random texture was assumed in the model based on texture analyses on hot rolled bcc steels found in the literature.

- For future simulations, it is suggested to use the formulation proposed by Vallellano et al. [51] for unequal grain sizes. N-R model used in this paper is strictly applicable to symmetric conditions including equal grain sizes. Beside,

- In fatigue tests with steel, multiple short cracks can grow simultaneously and form a main crack that propagates until failure. However, in this work, it was assumed that only a single crack initiates and grows until the end of initiation life, defined as a crack size of 10 grains. Therefore, no macrocrack growth and crack coalescence, which can occur in reality, were considered in the model, although many initiation spots can occur at the different ribs and only the worst combination leads to a dominating crack and failure.

### 4.6 Acknowledgements

We are grateful to Dr. Emmanuelle Courjault from the Computational Materials Laboratory (LSM) at EPFL for the EBSD analysis. We are also grateful also to Dr. Cayron from the Nanocharacterisation group of LITEN at CEA-Grenoble for the parent austenite reconstruction.
Bibliography


Fatigue behaviour prediction of steel reinforcement bars using an adapted Navarro and De Los Rios model

— Marina Rocha, Eugen Brühwiler, Alain Nussbaumer

**Abstract:** Fatigue cracks tend to initiate on the rebar surface and therefore, the surface conditions may control their fatigue behaviour. This study investigates the influence of surface microstructure and roughness dispersion on the scatter and fatigue life of hot rolled (HR)-cold worked (CW) and quenched and self-tempered (QST) rebars. The stochastic nature of the fatigue life is mainly affected by the scatter of short cracks in the crack initiation phase. A model adapted from Navarro and De Los Rios was developed to predict the crack initiation, including short crack growth, and long crack propagation phases. The stress concentration factor determined near the ribs was considered as a constant parameter. The model results were compared to experimental data from the literature.

**Keywords:** Steel reinforcement bars; Crack growth model; Surface conditions; Scatter; Fatigue behaviour prediction.
5.1 Introduction

Fatigue life of structural metallic components until failure can be split into initiation and propagation periods. Initiation is generally defined as the smallest crack that can be detected by non-destructive inspection technique. Propagation follows and in a cracked component, Paris’ law is usually applied for fatigue life assessment describing the growth rate from the detected initial-crack. In reinforced concrete elements, where crack detection is impracticable on the embedded steel reinforcement bars (rebars), S-N curve-method is used for fatigue life prediction. However, this method doesn’t provide any information on the presence, or not, of fatigue cracks. Paris’ law is only applied when a more detailed investigation is required with a long or conservative initial crack size being assumed in the calculations.

Fatigue strength of metallic materials may be controlled by the short crack growth behaviour as a consequence of the strength offered by the barriers, such as grain-phase boundaries, to the plastic slip. Short cracks can behave significantly different from long cracks predicted by Paris’ law [1]. These cracks may propagate during a large fraction of the component life and therefore, Paris’ law would fail to predict their fatigue life.

Fatigue cracks tend to initiate at stress raisers on the material surface. As a consequence, the conditions of the surface layer such as roughness and geometrical features is significant for the fatigue behaviour of the material. These stress raisers
are introduced in the fabrication process and they can be, in some cases, essential to a component perform its function.

Surface conditions affect primarily the crack initiation period, especially near the fatigue limit. Initiation, in this case, includes crack nucleation and short crack growth controlled by material barriers. Nucleation can be dependent on local surface irregularities which vary from specimen to specimen and can affect the duration of the crack initiation period. As a consequence, more scatter is found at high number of stress cycles [2].

The stress concentration on a surface irregularity (notch) is usually quantified in terms of stress concentration factor $K$. As $K$ increases, the stress required to initiate a crack at the notch decreases. It has been observed that the short crack growth rate at the notch root of steels, for example, can increase with the increase of $K$ values although these cracks can arrest or become non-propagating at the notch root after overcoming few barriers [3, 4]. This suggests that the initiation of the crack itself isn’t the key point in the fatigue behaviour of notched steels but rather the capacity of the (short) crack to propagate over successive microstructural barriers.

In this work, the fatigue behaviour of hot rolled (HR), cold worked (CW) and quenched and self-tempered (QST) rebars is investigated using an adapted Navarro and De Los Rios (N-R) model as given in [2]. The model includes the influence of the microstructure as well as the $K$ dispersion obtained from surface roughness analysis. The $K$ value determined for the rib geometry at the critical zone i.e., from where fatigue cracks usually initiate, is considered as a constant. The results are compared to experimental data for HR-CW and QST rebars.

### 5.2 Crack growth model

The algorithm used in this work is adapted from the N-R model described in [2] for plain specimens and represented in Fig. 5.1. The present model includes the roughness dispersion determined on the rebar surface as well as the long crack propagation phase to the N-R model [2] to investigate the fatigue behaviour of HR-CW and QST rebars.

As shown in Fig. 5.1, the crack initiation phase consists of a stochastic process which includes the dispersion inherent to:

- grain size variation;
- grain orientation ratio (for randomly oriented grains);
\textbf{Figure 5.1} – Flowchart of the adapted N-R model used in this work.
• different phases (ferrite-pearlite and martensite);

• stress concentration factors determined from surface roughness analysis.

In the long crack propagation phase, there is no dispersion of the microstructural features and therefore, it is assumed equal grain sizes and consequently equal increments of the plastic zone. The plastic zone size is negligible compared to the crack size.

5.2.1 Threshold for short crack growth

The short crack growth model for plain specimens [2] defines the condition to activate a plastic slip in terms of applied stress level and crack size; short cracks are unable to overcome microstructural barriers, such as grain-phase boundaries, at stress levels below the fatigue limit \( \Delta \tau_{FL} \). This interpretation is in agreement to what has been observed by [3, 5].

The applied stress \( \Delta \tau_{Li} \) required to propagate a short crack over \( i \) grains is given by [2]:

\[
\Delta \tau_{Li} = \Delta \tau_{FL} \frac{m_i}{m_1} \sqrt{\frac{d_i}{2c_i}}
\]

where \( \Delta \tau_{FL} \) is the fatigue limit of the plain specimen, \( d_i \) is the mean of the crack length in each grain, \( c_i \) is the position of the plastic zone and \( m_i / m_1 \) is the grain orientation ratio.

When irregularities (notches) are present on the specimen surface, the crack and its plastic zone growth are controlled by the resistance offered by the grain boundary as it occurs in plain specimens. However, the main difference in the short crack growth behaviour between plain and notched cases is the stress gradient related to the notch: the driving stress can vary significantly as the crack confronts each grain boundary. Depending on the applied stress level and the severity of the stress gradient, a short crack may grow over few grains and then stop as the stress level decreases. The crack propagates to the next grain only if the plastic slip is activated beyond the grain boundary.

The applied stress \( \Delta \tau_{Li}^{\text{notch}} \) required for the crack to overcome the \( i \)-th barrier in a notched specimen is given as [6]:

\[
\Delta \tau_{Li}^{\text{notch}} = \Delta \tau_{Li} K_f
\]

where \( K_f \) is the fatigue stress concentration factor. \( K_f \) values determined from sur-
face roughness analysis on QST rebar is discussed in Section 5.3.2.

5.2.2 \( f \) function

The crack propagation rate \( da/dN \) in the N-R model depends on the \( f \) function and it is given by:

\[
\frac{da}{dN} = f \phi \tag{5.3}
\]

where \( f \) represents the fraction of dislocations ahead of the crack tip that contributes to the crack growth process. \( N \) is the number of cycles and \( \phi \) is the crack tip plastic displacement. Since \( f \) varies as a function of the applied stress (it decreases as the applied stress level decreases) and the fatigue behaviour of rebars in this work is analysed at different stress levels, \( f \) functions were then proposed for ferrite, pearlite and martensite depending on the applied stress.

In the N-R model, \( f \) is constant at each stress level and is always obtained experimentally. In this work, three equations for the different phases were deduced on the experimental data with low Carbon steel under uniaxial loading as given in [7]. Depending on the fatigue limit of each phase, it was then assumed the same growth rate as found in the literature.

\( f \) functions determined for ferrite, pearlite and martensite are given in Eqs. 5.4, 5.5 and 5.6 respectively:

\[
f = 4.89 \times 10^{-16} \Delta \tau^{5.49} \tag{5.4}
\]

\[
f = 1.93 \times 10^{-20} \Delta \tau^{7.01} \tag{5.5}
\]

\[
f = 8.30 \times 10^{-20} \Delta \tau^{6.81} \tag{5.6}
\]

5.3 Surface roughness

Surface roughness is usually associated with the geometric topography of material surface. It depends greatly on the production technique where each fabrication process generates its own characteristic surface. Surface roughness manifests as a sequence of micronotches from where slip bands can emerge. The stress concentration arising from these micronotches may accelerate the fatigue crack nucleation
and lead to early short crack growth. In rebars, these micronotches are on the free surface and mainly concentrated near the ribs (see Fig. 5.2).

![Image of rebar axis direction](image)

**Figure 5.2** – Roughness on the surface of a QST rebar near the rib.

### 5.3.1 3D surface roughness profile

A photometric stereo technique applied to XL30-FEG Scanning Electron Microscopy (SEM) was used to reconstruct the 3D surface roughness profile of the QST rebar with diameter of 16 mm. An area of 1500 × 1500 μm, close to a transversal rib, was considered in the analysis. Four images were captured from the same area. The 3D surface reconstruction was then obtained by post-processing of all images based on photometric stereo technique. Fig. 5.3 shows the 3D surface roughness profile reconstructed by photometric stereo technique using SEM images.

The effect of the surface roughness on the fatigue strength of rebars can be obtained as a function of $R_a$, $R_y$ and $R_z$ parameters. They were obtained from the roughness profile height distribution ($z$) recorded over a length ($L$) and calculated as [8]:

$$R_a = \frac{1}{L} \int_0^L |z| \, dx$$  \hspace{1cm} (5.7)

where $R_a$ is the average deviation in surface height from the roughness profile mean line.

$$R_y = |z_{max} - z_{min}|$$  \hspace{1cm} (5.8)

$R_y$ is the maximum peak-to-valley height roughness over the evaluated length.

$$R_z = \frac{1}{5} \left( \sum_{r=1}^{5} (z_r)_{max} + \sum_{s=1}^{5} |(z_s)_{min}| \right)$$  \hspace{1cm} (5.9)

$R_z$ is the ten-point roughness i.e., the average height from the five highest peaks.
and five lowest valleys. Fig. 5.4 illustrates a surface roughness profile determined for the QST rebar over a length $L=1500 \mu m$. The dash line represents the mean of the roughness profile. The circles correspond to the five highest peaks and five lowest valleys considered in the $R_z$ calculations.

**Figure 5.3** – 3D profile of the surface roughness of the QST rebar obtained by photometric stereo technique.

**Figure 5.4** – Illustration of a surface roughness profile obtained for the QST rebar and the surface roughness parameters.
5.3.2 Fatigue stress concentration factor

The fatigue stress concentration factor $K_f$, imposed by the surface roughness and considered in the model, was calculated as [8]:

$$K_f = 1 + q \left[ n \left( \frac{R_d}{\bar{p}} \right) \left( \frac{R_y}{R_c} \right) - 1 \right]$$  \hspace{1cm} (5.10)

where $q$ is the notch sensitivity, $n=2$ for uniform tension stress and $\bar{p}$ is the effective profile valley radius. It represents the average radius obtained from the dominant profile valleys. The parameter $q$ is determined as:

$$q = \frac{1}{1 + y/\bar{p}}$$  \hspace{1cm} (5.11)

where $y$ is a material constant and defined as a function of the ultimate tensile strength $\sigma_u$ for steels [8]:

$$y = 0.025 \left( \frac{2070 \text{MPa}}{\sigma_u} \right)^{1.8}$$  \hspace{1cm} (5.12)

Since there are no studies in the literature on the surface roughness profile of rebars, the $K_f$ values determined in this work for the QST rebar were applied in the model for HR-CW rebars. The mean $\sigma_u$ of QST rebars [9] was then used in the calculations of $K_f$. For more accurate results, it is suggested to determine the roughness profile of HR-CW rebars and other QST rebars.

The $K_f$ values, calculated according to Eq. 5.10, were determined from the 3D surface roughness profile given in Fig. 5.3. The $K_f$ dispersion is shown in Fig. 5.5 with a mean value of 1.3.

The surface roughness of rebars is affected by the introduction of the ribs during the fabrication process; a higher surface roughness is observed near the transversal ribs (see Fig. 5.2). The material properties considered in the model are based on a literature review and they include implicitly unknown surface roughness $K_f$ greater than 1. The use of a mean $K_f$ value equal to 1.3 in the model could therefore underestimate the fatigue life predictions. For this reason, it is proposed to replace the mean $K_f = 1.3$ by 1, but keeping the scatter. The distribution of the $K_f$ values, given in Fig. 5.5, was approximated to a normal law with a mean value of 1 and a standard deviation of 0.1.
5.4 Stress concentration factor-rib geometry

Fatigue cracks in rebars usually initiate at the transversal rib base region (a zone between the weld toe radius and the rebar cylinder) [9, 10]. This critical region, illustrated in Fig. 5.6, is located along the ribs and it shows high stress concentration factors $K_t$ values on the rebar surface [11].

Figure 5.6 – Illustration of the critical zone along the ribs considered in this work.

The rib geometry details and the respective stress concentration factors are mostly unknown for the experimental data considered in this paper. However, these data, especially for QST rebars, consist mainly of specimens with similar rib patterns. Moreover, an average $K_t = 1.6$ can be expected on the zone along the transversal...
ribs [11] where fatigue cracks can initiate. For this reason, the effect of the rib geometry was considered by a constant $1/K_f$ factor multiplied by each stress range level in the model for HR-CW and QST rebars. A similar approach is given in [12].

### 5.5 Results and discussion

The influence of microstructure, surface roughness ($K_f$) and rib geometry ($K_t$) on the scatter and fatigue behaviour of HR-CW and QST rebars was analysed in this work. A total of 1000 simulations were run for different stress range levels. The material properties considered in this model are given in [2]. Ferrite-pearlite (F-P) and tempered martensite (TM) refers to the models for HR-CW and QST rebars, respectively.

The model included both crack initiation and propagation phases. The initiation phase is a stochastic process which includes the growth of microstructurally short cracks. Short cracks are allowed to grow to a maximum length of $2a$ equal to 10 grains from where the macro crack propagation phase starts [13] (see Fig. 5.1). A short crack propagates if $\Delta \tau_{\text{notch}}^m$ is greater than $\Delta \tau$. If this condition is not satisfied, the short crack stops to propagate and a run-out result is obtained in the model. Run-out result is, therefore, a non-propagating micro crack in an analysed region near the rib; if the short crack growth stops, it will no longer propagates. Since the simulations were run to utmost 100 million cycles, run-out results obtained at any number of cycles in the simulations represent run-outs at 100 million cycles as well.

The propagation phase in the model shows a macro crack growth where the influence of microstructural features such as grain-phase boundaries and grain orientation ratio is negligible. For this reason, similar grain sizes were considered with constant increments of the plastic zone $c_i = D_{\text{mean}}$. In fatigue testing of rebars with $d=16$ mm, when a surface crack length $2a$ of approximately 8 mm was detected, a rapid failure was in process [9]. Since rebars with $d \leq 16$ mm are also considered in the experimental data, the fatigue behaviour prediction would be overestimated for a crack length of 8 mm. For this reason, rebar failure was assumed, in all cases, for a surface crack length $2a = 6$ mm in the model.

Figs. 5.7 to 5.10 show the F-P and TM model results obtained for the crack initiation and propagation phases compared to experimental data. Experimental data is always represented in black and model results is represented in other colors. These data consist of HR-CW and QST rebars with diameter $d \leq 16$ mm [9, 14–17].
Figs. 5.7 and 5.9 show the model data points obtained for the crack initiation phase predicted for HR-CW and QST rebars, respectively. The green marks represent the number of cycles for a crack to overcome the first grain boundary at an applied stress range level. The blue marks correspond to the fatigue life of a short crack that propagated over 10 grains. Since the average grain size considered in the model for HR-CW and QST rebars is about 20 μm [2, 18], the crack length at the end of the initiation phase is approximately 0.2 mm.

The model results given in Figs. 5.7 to 5.10 show that a large fraction of the fatigue life of HR-CW and QST rebars is occupied by the crack initiation phase. This phase consists of the short crack growth to a maximum length of 0.2 mm and it covers approximately 80 (± 10) % and 60 (± 10) % near the fatigue limit of HR-CW and QST rebars, respectively. Paris’ law is, therefore, not appropriate for fatigue life predictions of rebars since it misses a significant part of the fatigue life prediction.

The difference between experimental and model data at high stress range levels (see Fig. 5.7) may be explained by the different pearlite area fractions that can be found in HR-CW rebars; their area fractions can vary between 15 and 55% while in the model a pearlite fraction of 53% was considered as given in [18]. Moreover, the higher scatter obtained in the initiation phase, especially for the crack to overcome the first grain boundary, is likely influenced by the presence of two different phases: the stress required to propagate in a pearlite grain, for example, is greater than the stress to overcome a (softer) ferrite grain [2].

The scatter trend obtained in the F-P and TM models, (see Figs. 5.7 to 5.10) was similar to the scatter obtained from experimental results, with an increase of scatter as the applied stress approaches the fatigue limit.

Figs. 5.11 and 5.12 shows the dispersion of the analysed parameters including only failure data points. To analyse the effect of a parameter in the model, the dispersion inherent to the other parameters was kept constant. The black points in Figs. 5.11 and 5.12 represent the influence of each parameter independently and the light grey points represent the dispersion of all the other parameters together. F-P and TM model results show that the influence of m_i/m, and grain size variation have similar effect on the scatter (see Figs. 5.11 and 5.12). Microstructure and roughness show approximately the same influence, near the fatigue limit, in both F-P and TM models; each parameter affects in about 50% the scatter at this region.
Figure 5.7 – Data points including the crack initiation phase (green marks: fatigue crack size = 1 grain; blue marks: fatigue crack size = 0.2 mm) in the F-P model and experimental data (black marks) of HR-CW rebars with diameter $D \leq 16$ mm [14,15].

### 5.6 Conclusion

The fatigue behaviour of HR-CW and QST rebars was predicted using an adapted N-R short crack growth model which includes stochastic crack initiation and linear propagation phases. The model takes into account the influence of the surface microstructure (grain orientation, grain size variation and phase) and roughness on the crack growth. The stress concentration factor from the rib geometry was considered as a constant parameter: it covers a critical region along the ribs which mostly show similar patterns. Based on the work presented in this paper, the following conclusions can be drawn:

- The main aspects influencing the fatigue behaviour of rebars, used as reference, were likely treated in this study using the F-P model and the TM models, since the scatter and fatigue behaviour predictions obtained from the models were similar to the experimental results. However, since fatigue test results from the literature include rebars with different rib patterns as well as residual stress at/near surface, the variation of these parameters may also influence their fatigue behaviour.
Figure 5.8 – Data points including the crack propagation phase (red marks) in the F-P model and experimental data (black marks) of HR-CW rebars with diameter \( D \leq 16 \) mm [14, 15].

- The fatigue life of HR-CW and QST rebars is largely occupied by the crack initiation phase. This phase consists of a crack which grows to a maximum length of 10 grains (or approximately 0.2 mm) in the model. Crack initiation covers, in general, 80 and 60% of the fatigue life near (but above) the fatigue limit of HR-CW and QST rebars, respectively. Paris’ law is, therefore, not appropriate to model fatigue of rebars since it misses most of the behaviour.

- The model shows that the influence of surface microstructure and roughness, on the scatter near the fatigue limit of HR-CW and QST rebars, is about 50% separately. TM model presents less scatter compared to F-P model and it shows that QST rebars have a better fatigue resistance and a higher fatigue strength than HR-CW rebars.
Figure 5.9 – Data points including the crack initiation phase (green marks: fatigue crack size = 1 grain; blue marks: fatigue crack size = 0.2 mm) in the TM model and experimental data (black marks) of QST rebar with diameter $D \leq 16$ mm [9, 16, 17].

Figure 5.10 – Data points including the crack propagation phase (red marks) in the TM model and experimental data (black marks) of QST rebar with diameter $D \leq 16$ mm [9, 16, 17].
Figure 5.11 – Influence of the - 1) Microstructure and roughness together (all); 2) Microstructure, including grain orientation ratio $m_1/m_2$, grain size variation, phases (ferrite-pearlite); 3) Each parameter of the microstructure separately and 4) Surface roughness - on the scatter obtained in the F-P model.
Figure 5.12 – Influence of the - 1) Microstructure and roughness together (all); 2) Microstructure, including grain orientation ratio \( m_i/m_1 \), grain size variation, phases (martensite); 3) Each parameter of the microstructure separately and 4) Surface roughness - on the scatter obtained in the TM model.
Bibliography


6.1 Introduction

This thesis applies concepts of the mechanics of materials and material engineering to define micro and macro structural aspects affecting the fatigue strength of different type of steel reinforcement bars (rebars). It provides advancements in the structural engineering methods used for fatigue life prediction. This research includes fatigue tests at very high number of stress cycles, micro-macro structural characterisation of rebars and an analytical model which predicts the main parameters influencing the fatigue behaviour of rebars.

6.2 Response to research questions

The main thesis contribution and findings to the research questions raised in Section 1.1 are concluded in this sub-chapter.

6.2.1 Fatigue testing

Experimental test data are used as reference to predict the fatigue behaviour of rebars in concrete structures. Axial tests of rebars alone have the advantage to be performed at very high number of stress cycles, i.e., greater than 10 million, at reduced time – and consequently costs – compared to fatigue tests with rebars embedded in concrete. However, axial fatigue tests are very sensitive to the grip arrangement which can lead to a rebar failure which isn’t characteristic of the proper fatigue searched behaviour. This research shows that conical grip is the most effective method preventing premature failure of the rebar inside the grip area. QST rebars mostly survived to a number of cycles exceeding 30 million at stress range levels of approximately 50% of their mean yield strength.
The method used for fatigue crack detection on these survival rebars, based on the application of the liquid penetrant testing (LPT) with an abrupt monitored frequency change (representative of fractured rebars), was ineffective. Survival rebars don’t show any abrupt frequency change; indeed, the frequency reaches a stabilization regime from about 1 million cycles on and kept nearly constant during the whole test. Moreover, LPT is a conservative method that can only detect rather large surface cracks of at least 5 mm length.

Surface imperfections were the determinant factor that led to the fracture of rebars, at similar stress ranges as applied for survival rebars. These imperfections are produced during the manufacturing process and located near the transversal ribs, in the zone of highest stress concentrations. Precisions regarding these surface imperfections are given in the next section.

6.2.2 Micro-macro structural characterisation

Conditions of the surface layer are most significant for the fatigue behaviour of rebars since fatigue cracks tend to initiate on the surface. This study provides new information on micro and macro surface aspects of QST rebars including experimental analyses of surface residual stresses and imperfections. The influence of geometric parameters on the stress concentration factors $K_t$ at the rebar surface was investigated by 3D Finite Element (FE) analyses. A parametric study using 3D FE models allowed to determine the influence of rib inclination and rebar size on the $K_t$ values of the critical zones. There were no references with similar studies of the surface residual stress, imperfections and 3D FE analysis of the geometric $K_t$ in the literature.

In this research, the microstructure of QST rebars was identified under Optical Microscopy and quantified for different rebar diameters. The analyses showed that the average grain size tends to increase with the increase of the rebar diameter.

Macroscopic residual stresses (type I) were experimentally determined on surface and subsurface of QST rebars by X-ray diffraction and Cut Compliance techniques. Residual stresses were found to be random tensile and compressive stresses with values no higher than 20% of their yield strength.

Surface imperfections and roughness, analysed under Scanning Electron Microscopy (SEM), are mainly located near the transversal ribs. The imperfections can be of different type:

- Marks near the transversal ribs;
• Marks with semi-circular shape near the transversal ribs;

• Cracks near the marks with some cracks perpendicular to the rebar axis. Cracks were also find perpendicular to the longitudinal ribs.

These imperfections are introduced in the rolling mill when the ribs are formed and they may also result from the water quenching process.

This research shows how the rib geometry and rebar diameter affects $K_i$ values on the rebar surface. The highest $K_i$ are located at the rib radius zone. The maximum $K_i$ values at this zone are mainly affected by the rib radius and height.

The highest $K_i$ values determined along the rib base increase with increasing rebar. The influence of the rebar diameter on these $K_i$ values is comparable to the changes in the rib radius and height. The maximum $K_i$ for QST rebars geometry with diameters of 10 and 26 mm are approximately 1.6 and 2.5, respectively. The effect of the rebar diameter on the size of the critical zone is less significant than the influence of rib radius and height.

The rib inclination is the main parameter affecting the position of the maximum $K_i$ and size of the critical zone along the rib. The critical zones (defined as zones where $K_i$ are at least 95% of the maximum values) are located anti-symmetrically along the rib. However, the position of the maximum $K_i$ tends to move to the rib centre and the size of the critical zone increases if the rib inclination becomes more and more perpendicular to the rebar axis.

6.2.3 Analytical model-part I

The scatter present in fatigue test results of rebars can be influenced by the sensitivity of the short crack growth to the microstructural features. This thesis proposes an adapted Navarro and De Los Rios (N-R) model within a Monte-Carlo framework to investigate the scatter above and near the fatigue limit on HR, CW and QST rebars. The N-R model, adapted in this work, incorporates the dispersion of the grain orientation ratio $m_i/m_{is}$, grain size variation and two phases (ferrite and pearlite) in the case of HR-CW rebars. A $m_i/m_{is}$ equation is developed to take into account crack closure effects and the variation of the grain orientations. Although N-R model has been extensively used for fatigue life prediction of metals, there are no references in the literature where the dispersion of microstructural features have been considered to analyse scatter in fatigue. Moreover, there are no references where a short crack growth model have been applied to study the scatter found in fatigue testing of rebars.
The developed approach shows that surface martensite contributes to reduce the scatter in fatigue tests with rebars compared to ferrite-pearlite microstructure found on the surface of HR-CW rebars. Surface microstructure affects to about 50% the scatter present in the fatigue tests for both HR-CW and QST rebars.

### 6.2.4 Analytical model-part II

The adapted N-R model in Section 6.2.3 is modified to include:

- the surface roughness effect in order to explain the difference in the scatter found in the analyses with the surface microstructure;
- a linear crack propagation phase where the effect of microstructural features is negligible in the crack growth.

The stress concentration factor from the rib geometry is considered as a constant parameter. This approach allows more realistic comparisons between model and experimental data. The model predicts the fatigue behaviour of HR-CW and QST rebars including fatigue crack initiation (cyclic slip, nucleation and short crack growth) and propagation (macro crack) phases.

This model shows fatigue behaviour predictions similar to test results for HR-CW and QST rebars. The main finding is that the fatigue life of rebars is largely occupied by the crack initiation phase during which the application of Paris’ law is infeasible.

This modelling demonstrates that QST rebars have a better fatigue resistance and a higher fatigue strength than HR-CW rebars. The standard S-N curves [1], used for safety verification, are nowadays classified according to the rebar size but there is no distinction between the fatigue strength of HR-CW and QST rebars. This approach is conservative and it is recommended to classify S-N curves according to the rebar type and size.

Fatigue cracks tend to initiate at the zone close to the ribs [2], [3], where the highest stress concentrations are present. The stress concentrations resulting from the rib geometry combined with the surface roughness near the ribs lead to a significant reduction on the fatigue strength of rebars. As a consequence, this finding suggests the use of plain rebars with surface martensite microstructure for fatigue relevant structures.
6.3 Future work

6.3.1 Fatigue crack detection

One of the greatest challenges in fatigue testing of rebars is the detection of cracks or microcracks, during or after the tests, in particular run-out tests. The methods used for crack detection are not efficient and therefore investigations are required in this area. One possible solution is to investigate the use of nanoparticles, in low surface tension solution, with further X-ray micro computed tomography analyses.

6.3.2 Grain orientation ratio

In this study, the equation proposed for the grain orientation ratio \( m_i/m_s \) was developed for randomly oriented grains. However, it is possible that HR, CW and QST rebars may show a preferential crystallographic orientation in the rolling direction. To provide more accurate microstructural information for the short crack growth modelling, the crystallographic texture of HR, CW and QST rebars could be determined by X-ray diffraction technique. Further research is needed on the effect of R-ratio in the fatigue behaviour of rebars. As given in [4], the \( m_i/m_s \) factor is affected by crack closure which partially governs the material resistance to fatigue damage. However, experimental data in the literature is still rather scarce for R-ratio greater than 0.2. The crack closure effect could be included in the \( m_i/m_s \) equation in the N-R model as proposed in this thesis.

6.3.3 Surface roughness

Further work is required in generating more information on the surface roughness profile near and between the ribs. This information could provide more accuracy on the effect of the surface roughness on the scatter present in fatigue tests and improve the fatigue behaviour prediction.

6.3.4 Fatigue behaviour modelling-other methods

For further modelling of fatigue behaviour, it is suggested to use the formulation proposed by Vallellano et al. [5] for non-symmetric conditions. Finite element (FE) modelling could be another promising method for a more detailed investigation on the stress field at each step of the short crack propagation, including the effect of local microresidual stresses, into and at the grain boundary. FE method has also the advantage to allow for the study of crack coalescence since the stress distribution of
the grains, of an analysed area, would not be uniform and micro cracks could start simultaneously at different grains.

The effect of variable amplitude loading could be studied by detailed modelling in parallel with lab tests to give a better understanding on the fatigue behaviour of rebars under the condition representative for structures subjected to fatigue.
Bibliography


Figure A.1 – Stress × Strain curve of a QST rebar analysed in this thesis.
Appendix A presents 3D Finite Element Models of the rebar geometry considered in the parametric study. The graphs given in this Appendix shows the evolution of the stress concentration factor, $K_t$, determined for the paths illustrated in Fig. B.1. Finite Element analyses were performed with Abaqus/CAE 6.12.

Figure B.1 – Paths (darker lines) where $K_t$ evolution was determined.
B.1 D1oh03r02

Figure B.2 – D1oh03r02 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.2 D10ho8ro2

Figure B.3 – D10ho8ro2 model: (a) 3D FE mesh (b) Top view (c) $K_i$ plot (d) Bottom view.
B.3  D10ho8r04

Figure B.4 – D10ho8r04 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.4  D1oh15r04

Figure B.5 – D1oh15r04 model: (a) 3D FE mesh (b) Top view (c) $K_I$ plot (d) Bottom view.
B.5 D1oh08

Figure B.6 – D1oh08 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.6 D16h03r02

Figure B.7 – D16h03r02 model: (a) 3D FE mesh (b) Top view (c) $K_i$ plot (d) Bottom view.
B.7  Dw16h08r02

Figure B.8 – Dw16h08r02 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.8  D16ho8ro4

Figure B.9 – D16ho8ro4 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.9 D16h08r08

Figure B.10 – D16h08r08 model: (a) 3D FE mesh (b) Top view (c) $K_i$ plot (d) Bottom view.
Figure B.11 – D16ho8 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.11 D16ho8r04c04

Figure B.12 - D16ho8r04c04 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
Figure B.13 – D16ho8r04c12 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.13 D26h03r02

Figure B.14 – D26h03r02 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.14  D26ho8ro2

![3D FE mesh](image1)

(a)

![Top view](image2)

(b)

![K plot](image3)

(c)

![Bottom view](image4)

(d)

Figure B.15 – D26ho8ro2 model: (a) 3D FE mesh (b) Top view (c) $K_i$ plot (d) Bottom view.
Figure B.16 – D26ho8ro4 model: (a) 3D FE mesh (b) Top view (c) $K_i$ plot (d) Bottom view.
Figure B.17 – D26h15r04 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
B.17  D26h15r08

Figure B.18 – D26h15r08 model: (a) 3D FE mesh (b) Top view (c) $K_t$ plot (d) Bottom view.
Appendix C

Model

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Appendix B presents the algorithm developed for the analytical model used in the analyses given in Chapters 4 and 5.

Figure C.1 - Model files: (a) file containing the input function to run an experiment; (b) the process will occur simultaneously in parallel on each core; (c) file containing all the function for the Voronoi structure and the propagation algorithm; (d) two files containing a class with properties for each of the two steels: TM & FP.
C.1 File experiment.m

% experiment.m file
% this is the main input function
% Examples:
% experiment(10,400,[180:200],1);
% experiment(10,400,linspace(180,200,1),1);

function matrix = experiment( N, points, applied_stress, is_martensite )

matlabpool local;

number_of_tests = N*numel(applied_stress);
matrix = zeros(8,number_of_tests);

value_time1 = clock;

value_ten = applied_stress(1);
parfor i = 1:3*3
    mainNR( {value_ten,points}, is_martensite );
end

value_time2 = clock;
print_estimation_time(number_of_tests,value_time1,value_time2);

tic

% parallel for loop (each step is an experiment)
parfor i = 1:number_of_tests
    current_stress = return_stress(applied_stress,i,N);
    [mem,~] = mainNR( {current_stress,points}, is_martensite );
    matrix(:,i) = [current_stress rot90(mem)];
end

tic

figure;
scatter(log10(matrix(2,:)),matrix(1,:),20);

figure;
plot_runouts_by_applied_stress
B=matrix(1:2,:);
B=(B==100000000);
get_N_values
Ns=matrix(B);
get_S_values
ml=B(2,:);
m2=zeros(7,length(B(2,:)));
matrix_select=cat(1,ml,m2);
Ss=matrix( matrix_select==1 ) ;
```matlab
%plot S-N curve
hist(cat(1,rot90(Ss)));
toc
matrix = rot90(matrix);
matlabpool close;

function result = return_stress(app,i,N)
    result = app(ceil(i/N));
end

function print_estimation_time(number_of_tests,value_time1,value_time2)
    init = value_time1(5)*60 + value_time1(6);
    stop = value_time2(5)*60 + value_time2(6);
    time_int = (stop - init)/(3*3);
    total = floor(time_int*number_of_tests);
    value_time3 = value_time2;
    value_time3(4) = floor(total/3600)+value_time3(4);
    value_time3(5) = floor(mod(total,3600)/60+value_time3(5));
    value_time3(6) = floor(mod(mod(total,3600),60)+value_time3(6));

    if value_time3(6) >= 60
        value_time3(5) = value_time3(5) + floor(value_time3(6)/60);
        value_time3(6) = mod(value_time3(6),60);
    end

    if value_time3(5) >= 60
        value_time3(4) = value_time3(4) + floor(value_time3(5)/60);
        value_time3(5) = mod(value_time3(5),60);
    end

    if value_time3(4) >= 24
        value_time3(4) = mod(value_time3(4),24);
    end

disp(['-> It will take ',num2str(total), ' seconds, for ...
     ',num2str(number_of_tests),' essays.']);
disp(['-> from ',num2str(value_time2(4)), ':', ...
     num2str(value_time3(4)), ':', num2str(floor(value_time2(6))), ... 
     ' to ', num2str(value_time3(4)), ':', num2str(value_time3(5)), ...
     ':', num2str(floor(value_time3(6)))]);
end
```

The text represents a MATLAB code snippet that includes plotting an S-N curve, some matrix manipulations, and a function to calculate the time taken for processing a certain number of essays.
C.2 File mainNR.m

% mainNR.m file
% this is the main input function
% Examples:
% [NumberCycles, Structure] = mainNR([49, 1000]);
% and to use an already made "Structure":
% [NumberCycles, Structure] = mainNR(Structure);

function [result_navarro, structure_data] = mainNR( varagin, ...
   is_martensite )

%constants
global calculus_error;
global print_slipbands_during_propagation;
global plot_result;
global load_orientation;
global global_dmean;
global number_of_grains_to_break;
global propagation_model;
global scale_structure;
result_navarro = zeros(7,1);
load_orientation = 0;
calculus_error = 1.0e-10;
plot_result = false;
print_slipbands_during_propagation = false;
number_of_grains_to_break = 10;
global_dmean = [0 0];
scale_structure = 1;
linear_propagation=true;
second_grain_limit=300;
experiment_statistics
statistics_matrix=ones(2,15,4);
	tic

if length(varagin) == 2
   applied_stress = varagin{1}; point_number = varagin{2};
   %create the grain structure
   [V, C, points, sb, maxgrainx, maxgrainy, maxgrain, maxfirstsb] = ...
      create_structure(point_number);
elseif length(varagin) == 11
   applied_stress = varagin{1}; point_number = varagin{2}; V = ...
   varagin{3}; C = varagin{4}; points = varagin{5}; sb = ...
   varagin{6}; maxgrainx = varagin{7}; maxgrainy = varagin{8}; ...
maxgrain = varargin{9}; maxfirstsb = varargin{10}; global_dmean ...
= varargin{11};
else
  return;
end
if is_martensite==1
  propagation_model = classTM(applied_stress, load_orientation);
else
  propagation_model = classFP(applied_stress, load_orientation);
end
PL_Structure = containers.Map({ -1, 1}, {struct('grain_id', maxgrain, ...
'slipband_id', maxfirstsb, 'init_point', points(maxgrain, :), ...
'end_point', [maxgrainx(1) maxgrainy(1)], 'length', ...
distance_segment(points(maxgrain, 1), points(maxgrain, 2), ...
maxgrainx(1), maxgrainy(1)), 'total_length', ...
distance_segment(points(maxgrain, 1), points(maxgrain, 2), ...
maxgrainx(1), maxgrainy(1)), 'angle', ...
get_angle_with_coordinates(points(maxgrain, 1), points(maxgrain, ...
2), maxgrainx(1), maxgrainy(1)), 'is_border', false), ...
struct('grain_id', maxgrain, 'slipband_id', maxfirstsb, ...
'init_point', points(maxgrain, :), 'end_point', [maxgrainx(2) ...
maxgrainy(2)], 'length', distance_segment(points(maxgrain, 1), ...
points(maxgrain, 2), maxgrainx(2), maxgrainy(2)), 'total_length', ...
distance_segment(points(maxgrain, 1), points(maxgrain, 2), ...
maxgrainx(2), maxgrainy(2)), 'angle', ...
get_angle_with_coordinates(points(maxgrain, 1), points(maxgrain, ...
2), maxgrainx(2), maxgrainy(2)), 'is_border', false});
CrackStructure = containers.Map({0}, {struct('c', 0, 'dmean', 0, ...
'nc', 0, 'ac', 0, 'ns', 0, 'dN', 0, 'N', 0, 'is_blocked', false});
if plot_result
  if length(get(0, 'children'))>1
    figure(2);
    clf;
    figure(1);
    clf;
  end
  plot_structure(V, C, points);
  %plot initial point
  plot(PL_Structure(1).init_point(1), PL_Structure(1).init_point(2), ...
'*', 'Color', 'black', 'linewidth', 4);% 'LineSmoothing', 'on');
  %plot slipbands(V, C, points, sb);
end
side_ok = [true true];
current_a = [0 0];
current_plastified_grain_step = [0 0];
lastminDaDN = - 1;
d_mean_stop = global_dmean(1, 1);
while sum(current_a) < number_of_grains_to_break*d_mean_step && ...
{side_ok(1, 1) == true || side_ok(1, 2) == true)
    if (Crack_Structure(current_plastified_grain_step(1, 1)).N <= ...
        Crack_Structure(current_plastified_grain_step(1, 2)).N && ...
        (side_ok(1, 1) == true && side_ok(1, 2) == true) || ...
        (side_ok(1, 1) == true && side_ok(1, 2) == false)
        index = 1;
        current_plastified_grain_step(1, index) = ...
        current_plastified_grain_step(1, index)+1;
    else
        index = 2;
        current_plastified_grain_step(1, index) = ...
        current_plastified_grain_step(1, index)-1;
    end
    if plot_result
        plot_plastic_zone(current_plastified_grain_step(1, index), ...
        PL_Structure);
    end
    PL_Structure = preferential_line(V, C, sb, PL_Structure, ...
    current_plastified_grain_step(1, index));
    [propagation_model, Crack_Structure, side_ok_res, matrix_disp] = ...
    propagation_model.is_propagating(current_plastified_grain_step(1, ...
    index), current_plastified_grain_step(1, mod(index, 2)+1), ...
    side_ok, Crack_Structure, PL_Structure);
%statistics
    statistics_matrix(index, abs(current_plastified_grain_step(1, ...
    index)), :) = matrix_disp;
    if plot_result
        disp(['PLASTIC ZONE: ', ...
            num2str(current_plastified_grain_step(1, index))]);
        disp(PL_Structure(current_plastified_grain_step(1, index)));
        disp(['CRACK INCREMENT: ', ...
            num2str(current_plastified_grain_step(1, index))]);
        disp(Crack_Structure(current_plastified_grain_step(1, index)));
    end
%When dN<=0, don't compute the dadN function
    if plot_result && side_ok(1, index) == true && ...
        Crack_Structure(current_plastified_grain_step(1, index)).dN<=0
        side_ok(1, index) = false;
        if index == 1;
            current_plastified_grain_step(1, index) = ...
            current_plastified_grain_step(1, index)-1;
        else
            current_plastified_grain_step(1, index) = ...
            current_plastified_grain_step(1, index)+1;
if ((Crack Structure(current plastified grain step(1, 1)).N <= ...)
Crack Structure(current plastified grain step(1, 2)).N ))
(side_ok(1, 1) == true && side_ok(1, 2) == true) || ...
(side_ok(1, 1) == true && side_ok(1, 2) == true)
current a(1, 1) = ...
Crack Structure(current plastified grain step(1, 1)).ac;
end
if ((Crack Structure(current plastified grain step(1, 2)).N <= ...)
Crack Structure(current plastified grain step(1, 1)).N ))
(side_ok(1, 1) == true && side_ok(1, 2) == true) || ...
(side_ok(1, 2) == true && side_ok(1, 1) == true)
current a(1, 2) = ...
Crack Structure(current plastified grain step(1, 2)).ac;
end
if plot_result && abs(current plastified grain step(1, 1)) > 0 && ...
abs(current plastified grain step(1, 2)) > 0
lastminDaDN = plotdadN(current plastified grain step, ...)
Crack Structure, PL Structure, lastminDaDN);
end
update here for the next step only (when the fracture stops or ...
arrives at the border
side_ok = side ok res;
if plot_result && side ok(1, index) == false
disp(['The crack stops at step ', ...
um2str(current plastified grain step(1, index))]);
end
if PL Structure(current plastified grain step(1, index)).is_border ...
== true
side ok(1, index) = false;
if plot_result
disp(['The crack arrives at the border at step ', ...
um2str(current plastified grain step(1, index))]);
end
end
if plot_result && abs(current plastified grain step(1, 1)) > 0 && ...
abs(current plastified grain step(1, 2)) > 0
d mean stop = (abs(current plastified grain step(1, ...1)) * Crack Structure(current plastified grain step(1, ...1)).d mean+abs(current plastified grain step(1, ...2)) * Crack Structure(current plastified grain step(1, ...2)).d mean)/(abs(current plastified grain step(1, ...1)) + abs(current plastified grain step(1, 2)));
end
store the value when the first grain breaks
if abs(current plastified grain step(1, 1)) == 1 && ...
abs(current_plastified_grain_step(1, 2))==1
result_navarro(2) = ...
max([Crack_Structure(current_plastified_grain_step(1, ...
1)).dN Crack_Structure(current_plastified_grain_step(1, ...
2)).dN]);

end
end

if PL_Structure(current_plastified_grain_step(1, 1)).is_border
side_ok(1, 1) = true;
elseif PL_Structure(current_plastified_grain_step(1, 2)).is_border
side_ok(1, 2) = true;
end

Ntotal = max([Crack_Structure(current_plastified_grain_step(1, 1)).N ...
Crack_Structure(current_plastified_grain_step(1, 2)).N]);

if side_ok(1, 1) == true || side_ok(1, 2) == true
if plot_result:
disp(['Propagation part: it breaks at N\text{_{propa}} = ', ...
num2str(Ntotal, '1.0\times10^5\n')]);
end
else
Ntotal = 10^8;
if plot_result:
disp(['Propagation part: the cracks stops before N = ', ...
num2str(Ntotal, '1.0\times10^5\n')]);
end
endif

%statistics
result_navarro(1) = Ntotal;
result_navarro(3) = Ntotal;
if (Crack_Structure(current_plastified_grain_step(1, ...
1)).N>Crack_Structure(current_plastified_grain_step(1, 2)).N)
stat_mat=statistics_matrix(1,1:abs(current_plastified_grain_step(1, ...
1)),:);
stat_mat=reshape(stat_mat,[abs(current_plastified_grain_step(1, ...
1)) 4]);
else
stat_mat=statistics_matrix(2,1:abs(current_plastified_grain_step(1, ...
2)),:);
stat_mat=reshape(stat_mat,[abs(current_plastified_grain_step(1, ...
2)) 4]);
end
result_navarro(4)=sum(stat_mat(:,1));
result_navarro(5)=sum(stat_mat(:,2));
result_navarro(6)=sum(stat_mat(:,3));
result_navarro(7)=sum(stat_mat(:,4));

structure_data = [applied_stress, point_number, V, C, points, sb, ...
maxgrainx, maxgrainy, maxgrain, maxfirstsb, global_dmean];
% linear propagation

if linear_propagation==true & & (side_ok(1, 1) == true || side_ok(1, 2) == true)
    c=[Crack_Structure(current_plastified_grain_step(1, 1)).c ... 
       Crack_Structure(current_plastified_grain_step(1, 2)).c];
    nc=[Crack_Structure(current_plastified_grain_step(1, 1)).nc ... 
       Crack_Structure(current_plastified_grain_step(1, 2)).nc];
    ns=[0 0];
    dN=[0 0];
    while abs(current_plastified_grain_step(1, ... 
        1))+abs(current_plastified_grain_step(1, ... 
        2))<second_grain_limit & & (side_ok(1, 1) == true || side_ok(1, ... 
        2) == true)
        for t=[-1 1]
            index_side = 1+(1-sign(t))/2;
            if side_ok(1, index_side) == true
                ns(index_side)=c(index_side)/(c(index_side) + ... 
                    d_mean_stop)*nc(index_side);
                c(index_side)=c(index_side)+d_mean_stop;
                current_plastified_grain_step(1, index_side) = ... 
                current_plastified_grain_step(1, index_side)+t;
                [dnRes,ncRes] = propagation_model.compute_long_crack( ... 
                    current_plastified_grain_step(1, index_side ), c( ... 
                    index_side ), d_mean_stop,ns(index_side) );
                dN(index_side)=dN(index_side)+dnRes;
                nc(index_side)=ncRes;
            end
            if plot_result
                disp(['CRACK INCREMENT: ', ... 
                    num2str(current_plastified_grain_step(1, ... 
                    index_side))]);
            end
        end
    end
    result.navarro(1)=result.navarro(1)+max(dN);
    if plot_result
        disp(['Linear part: N linear = ', num2str(max(dN), '%10.5e\n')]);
    end
end

function plot_plastic_zone(k, PL_Structure)
    if ~isempty(get(0, 'children'))
        figure(1);
if abs(k) == 1
    plot([PL_Structure(k).init_point(1) PL_Structure(k).end_point(1)], ...%
        [PL_Structure(k).init_point(2) PL_Structure(k).end_point(2)], ...%
        'Color', [0 0 0], 'linewidth', 4);% 'LineSmoothing', 'on');
else
    plot([PL_Structure(-1).end_point(1) PL_Structure(-1).end_point(1)], ...%
        [PL_Structure(-1).end_point(2) PL_Structure(-1).end_point(2)], ...%
        'Color', [0 0 0], 'linewidth', 4);% 'LineSmoothing', 'on');
end

function plot_crack(a_p, a_m, PL_Structure)
if ~isempty(get(0, 'children'))
    figure(1);
else
    figure('Name', 'Grain structure');
end
for i = [1 -1]
    current_length = 0;
    if i == 1
        a = a_p;
        grain_id = 1;
    else
        a = a_m;
        grain_id = -1;
    end
    if numel(a)>1
        while current_length<a(2)
            new_length = current_length+PL_Structure(grain_id).length;
            if a(1)<= new_length
                prop_coef = (a(1) - ...
                                current_length)/PL_Structure(grain_id).length;
                points1 = ([ PL_Structure(grain_id).end_point(1) ...%
                                PL_Structure(grain_id).init_point(1) ...%
                        )*prop_coef + ...%
                        PL_Structure(grain_id).init_point(1) ( ...%
                        PL_Structure(grain_id).end_point(2) - ...%
                        PL_Structure(grain_id).init_point(2) ...%
                        )*prop_coef + ...%
                        PL_Structure(grain_id).init_point(2));
            else
                points1 = PL_Structure(grain_id).init_point;
            end
            if a(2)>= current_length & a(2)<new_length
                prop_coef = (a(2) - ...
current_length)/PL_Structure(grain_id).length;
points2 = [( PL_Structure(grain_id).end_point(1) - ...
PL_Structure(grain_id).init_point(1) ... )*prop_coef + ...
PL_Structure(grain_id).init_point(1) ( ...
PL_Structure(grain_id).end_point(2) - ...
PL_Structure(grain_id).init_point(2) ... )*prop_coef + ...
PL_Structure(grain_id).init_point(2)];
else
points2 = PL_Structure(grain_id).end_point;
end
if sign(i) == - 1, color = 'red'; else color = 'blue'; ... end;
plot([points1(1) points2(1)], [points1(2) points2(2)], ...
'Color', 'white');
end
current_length = current_length+PL_Structure(grain_id).length;
grain_id = grain_id+1*i;
end
e}
end
else
points2 = PL_Structure(grain_id).end_point;
end
end
end
end
end

function mindaDN = plotdadN(steps, Crack_Structure, PL_Structure, mindaDN)

global propagation_model;

if length(get(0, 'children'))>1
figure(2);
else
figure('Name', 'da/dN');
end

offset_alone = 0;%when the loop is only k

minX = [0 0];
maxX = [0 0];

if Crack_Structure(steps(1, 1)).N >= Crack_Structure(steps(1, 2)).N
k = steps(1, 1);l = steps(1, 2);
else
k = steps(1, 2);l = steps(1, 1);
end

%lap length computations
if Crack_Structure(k - sign(k)).N > Crack_Structure(l).N
loop = k;
%when a crack stops, fill the gap
if Crack_Structure(k - 2*sign(k)).N <= Crack_Structure(l).N
[propagation_model, minX(l)] = ... propagation_model.crack_position_by_dN(Crack_Structure(l).N ...
- Crack\_Structure(k - 2*\text{sign}(k)).\text{N}, Crack\_Structure(k - \text{sign}(k)).\text{ns}, Crack\_Structure(k - \text{sign}(k)).\text{c}, k - \text{sign}(k), ...
l, Crack\_Structure, PL\_Structure);

```matlab
else
    \text{minX}(1) = Crack\_Structure(k - \text{sign}(k)).\text{ac};
end
maxX(1) = Crack\_Structure(k).\text{ac};
offset\_alone = Crack\_Structure(l).\text{ac};
else
loop = [k l];
    if Crack\_Structure(k - \text{sign}(k)).\text{N} = Crack\_Structure(l - \text{sign}(l)).\text{N}
        \text{minX}(1) = Crack\_Structure(k - \text{sign}(k)).\text{ac};
        maxX(2) = Crack\_Structure(l).\text{ac};
    end
    \text{compute minX for l, when N = N(k - 1)}
    [\text{propagation\_model, minX}(2)] = ...
        \text{propagation\_model.crack\_position\_by\_dN(Crack\_Structure(k - ...}
            \text{sign(k)).\text{N}, Crack\_Structure(l - \text{sign}(l)).\text{N}, ...}
        \text{Crack\_Structure(l).\text{ns}, Crack\_Structure(l).\text{c}, l, k - ...}
            \text{sign(k), Crack\_Structure, PL\_Structure});
    \text{compute maxX for k, when N = N(l)}
    [\text{propagation\_model, maxX}(1)] = ...
        \text{propagation\_model.crack\_position\_by\_dN(Crack\_Structure(l).\text{N} ...}
            \text{- Crack\_Structure(k - \text{sign}(k)).\text{N}, Crack\_Structure(k).\text{ns}, ...}
        \text{Crack\_Structure(k).\text{c}, k, l, Crack\_Structure, PL\_Structure});
end all l - N segments are inside the k - N segment
```

```matlab
\text{minX}(2) = Crack\_Structure(l - \text{sign}(l)).\text{ac};
maxX(2) = Crack\_Structure(l).\text{ac};
[\text{propagation\_model, minX}(1)] = ...
    \text{propagation\_model.crack\_position\_by\_dN(Crack\_Structure(l).\text{N} ...}
        \text{- Crack\_Structure(k - \text{sign}(k)).\text{N}, Crack\_Structure(k).\text{ns}, ...}
    \text{Crack\_Structure(k).\text{c}, k, l, Crack\_Structure, PL\_Structure});
[\text{propagation\_model, maxX}(1)] = ...
    \text{propagation\_model.crack\_position\_by\_dN(Crack\_Structure(l).\text{N} ...}
        \text{- Crack\_Structure(k - \text{sign}(k)).\text{N}, Crack\_Structure(k).\text{ns}, ...}
    \text{Crack\_Structure(k).\text{c}, k, l, Crack\_Structure, PL\_Structure});
end
end
```

numb\_int\_points = 10;
X = [1: numb\_int\_points];
Y = zeros(numb\_int\_points, 1);
if size(loop) == 1
    if \text{sign}(k) == 1, \text{color} = 'blue'; else \text{color} = 'red'; end;
else
    if \text{sign}(k) == 1, \text{color} = 'red'; else \text{color} = 'blue'; end;
end

factor\_To\_Meter\_By\_Cycle = 10^-3; % because unity in Mpa (N/mm2)
if sum(maxX) == 0
    for p = 1:numel(loop)
```
i = loop(p);
step = (maxX(p) - minX(p))/(numb_int_points - 1);
for j = 1: numb_int_points
  a = minX(p) + (j - 1)*step;
  [propagation_model, newdadn] = propagation_model.dadN(a, ...
    a/Crack_Structure(i).c, k, 1, Crack_Structure, ...
    PL_Structure);
  Y(j) = Y(j) + newdadn;
end
plot(sum(minX)+(sum(maxX) - sum(minX))/(numb_int_points - 1)*(X - ... 1) +offset_alone, (Y*factorToMeterByCycle), 'black');
end
if mindaDN>0
  plot([sum(minX)+offset_alone sum(minX)+offset_alone], ([mindaDN ... Y(1)]*factorToMeterByCycle), 'black');
end
%grain boundaries
hold on;
X2 = {{[Crack_Structure(k).c+Crack_Structure(l).c ...
  Crack_Structure(k).c+Crack_Structure(l).c])};
Y2 = {[min(Y) max(Y)]*factorToMeterByCycle};
plot(X2, Y2, '--', 'Color', color);
%plot crack
if sign(k) == 1
  plot_crack([minX(1) maxX(1)], [minX(2) maxX(2)], PL_Structure);
else
  plot_crack([minX(2) maxX(2)], [minX(1) maxX(1)], PL_Structure);
end
mindaDN = min(Y);
end
%create the voronoi structure
function [V, C, points, sb, first_slip_band_x, first_slip_band_y, ...
  init_grain_id, maxfirstsb_angle_index] = ...
  create_structure(point_number)
points = zeros(point_number, 2);
maxfirstsb_angle_index = -1;
%here to change the grain strucure
for a = 1:point_number
  %pointx(a) = random('norm', 0.5, 0.2);
  %random
  points(a, 1) = random('unif', -0.5, 1.5);
  points(a, 2) = random('unif', -0.5, 1.5);
  %hexa
  %offset = mod(floor(a/sqrt(point_number)), 2)/sqrt(point_number);
end
end

global scale_structure;
*scale_structure = 10^-3;
points = points*scale_structure;
[V, C] = voronoin(points);

%sb = slip bands
%psb = first slip band
number_of_slip_band = 3;
sb = zeros(point_number, number_of_slip_band);
for a = 1:point_number
    %psb = 0;
    psb = random('unif', 0.0, pi/number_of_slip_band);
    for k = 1:number_of_slip_band
        sb(a, k) = psb+(k - 1)*pi/number_of_slip_band;
    end
end
maxlc = 0;
init_grain_id = 0;
first_slip_band_x = 0;
first_slip_band_y = 0;
for grain = 1:point_number
    if points(grain, 1)>0 && points(grain, 2)>0 && points(grain, ... 1)<scale_structure && points(grain, 2)<scale_structure
        [grainx, grainy, sb_index, sb_angle] = find_slip_band( V, C, ... grain, sb(grain, :), points(grain, 1), points(grain, 2), ... []], 0 );
        if numel(grainx)>1
            sb_length = distance_segment(grainx(1), grainy(1), ... grainx(2), grainy(2));
```matlab
%compute lc
lc = abs(sb_length*cos(2*(pi/4 - (sb(grain, sb_index)))));

if lc>maxlc
    maxlc = lc;
    init_grain_id = grain;
    first_slip_band_x = grainx;
    first_slip_band_y = graniy;
    maxfirstsb_angle_index = sb_index;
end
end

%compute the global grain mean size here
global global_dmean;
global_dmean(1, 1) = global_dmean(1, 1)/global_dmean(1, 2);
end

%plot the voronoi structure
function plot_structure(V, C, points)
%global scale_structure;
scale_structure = 1;

if length(get(0, 'children'))>1
    figure(1);
else
    figure('Name', 'Grain structure');
end

for a = 1:length(points(:, 1))
    hold on;
    if (a>1)
        %plot(pointx(a - 1:a), pointy(a - 1:a));
    end
    %text(points(a, 1), points(a, 2), num2str(a), 'FontSize', 8, ...
         'Color', 'black');
    end
hold on;
%plot(points(:, 1), points(:, 2), '*', 'Color', 'k', 'MarkerSize', 2);
%plot grain: if convhull raises an error, other method to draw the points.
try
    for k = 1:length(C)
        if all(C{k}== 1)
            VertCell = V(C{k}, :);
            KVert = convhulln(VertCell);
            patch('Vertices', VertCell, 'Faces', KVert, ...
end

end

catch errN
    for j = 1:length(C)
        b = C{j, :};
        number_vert = size(b);
        for i = 1:number_vert(2)
            hold on;
            if (i>1)
                plot([V(b(i-1), 1) V(b(i), 1)], [V(b(i-1), 2) V(b(i), ... 2)], 'Color', [0 0 0], 'LineWidth', 0.1);
            else
                plot([V(b(1), 1) V(b(number_vert(2), 1))], [V(b(1), 2) ... V(b(number_vert(2), 2))], 'Color', [0 0 0], ... 'LineWidth', 0.1);
            end
        end
    end
end

axis equal;
axis([-0.5 1.5 -0.5 1.5]*scale_structure);

%plot the slipbands
function plot_slipbands(V, C, points, sb)
    for grain = 1:length(points(:, 1))
        find_slipband( V, C, grain, sb(grain,:), points(grain, 1), ... points(grain, 2), [], true);
    end
end

%find the equation of the line
function [A, B] = line_equation( X, Y )
    A = (Y(1) - Y(2))/(X(1) - X(2));
    B = Y(2) - A*X(2);
end

%give the length of the segment
function length_seg = distance_segment(x1, y1, x2, y2)
    length_seg = sqrt((x1 - x2)^2 + (y1 - y2)^2);
end

%give the next grain
function boundary_grain = find_next_grain( V, C, grain, x_inter, ... y_inter )
global calculus_error;
boundary_grain = 0;

test = C{grain, :};
tamanho = size(test);

%loop on vertices
for i = 1:tamanho(2)
    x1 = V(test(i), 1);
    y1 = V(test(i), 2);
    first_point = test(i);
    if i == tamanho(2)
        x2 = V(test(1), 1);
        y2 = V(test(1), 2);
        second_point = test(1);
    else
        x2 = V(test(i+1), 1);
        y2 = V(test(i+1), 2);
        second_point = test(i+1);
    end

    %check if the line is/not vertical
    if abs(x1 - x2) < calculus_error
        erro = abs(x_inter - x1);
    else
        [A, B] = line_equation([x1 x2], [y1 y2]);
        erro = abs(A*x_inter+B - y_inter);
    end

    %disp(['erro ', num2str(erro)]);
    if erro < calculus_error
        for j = 1:length(C)
            tem_first_point = 0;
            tem_second_point = 0;
            vertices_list = C{j, :};
            tamanhoc = size(vertices_list);
            for k = 1:tamanhoc(2)
                if vertices_list(k) == first_point
                    tem_first_point = 1;
                elseif vertices_list(k) == second_point
                    tem_second_point = 1;
                end
            end
            if tem_first_point == 1 && tem_second_point == 1 && grain ~== j
                boundary_grain = j;
            end
        end
    end
end
function [maxgrainx, maxgrainy, slip_band_index, ...,
     slip_band_oriented_angle] = find_slip_band( V, C, grain, anglesb, ...,
     x_init, y_init, first_step_sb, print)

maxlc = 0;
minlc = pi;
slip_band_index = 0;
slip_band_oriented_angle = 0;
global calculus_error;
global load_orientation;
global global_dmean;
global scale_structure;
scale_structure = 1;
maxgrainx = 0;
maxgrainy = 0;
b = C{grain, :};
tamanho = size(b);
tamanho = tamanho(2);
vgx = zeros(tamanho+1, 1);
vgy = zeros(tamanho+1, 1);

for i = 1:tamanho
    vgx(i) = V(b(i), 1);
    vgy(i) = V(b(i), 2);
end

% close the grain
vgx(tamanho+1) = V(b(1), 1);
vgy(tamanho+1) = V(b(1), 2);

for sbg = 1:length(anglesb)
    linex = [cos(anglesb(sbg))+x_init cos(anglesb(sbg)+pi)+x_init];
    liney = [sin(anglesb(sbg))+y_init sin(anglesb(sbg)+pi)+y_init];
    [xi, yi] = polyxpoly(vgx, vgy, linex, liney);

    if length(xi)>1 && max(abs(max(vgx - 0.5*scale_structure)), ...
                       abs(min(vgx - 0.5*scale_structure))) <= 1.0*scale_structure && ...
                   max(abs(max(vgy - 0.5*scale_structure)), abs(min(vgy - ...)
                        0.5*scale_structure))) <= 1.0*scale_structure & assegurar que ...
todas as linhas formam um polígono fechado

if abs(xi(2) - x_init) < calculus_error
    disp(['need to invert the matrix init (' ...  
        num2str(x_init), ', ', num2str(y_init), ') and ... ' ...  
        num2str(xi(2))]);
    tempx = [xi(2) xi(1)];
    tempy = [yi(2) yi(1)];
    xi = tempx;
    yi = tempy;
end

if print == true
    if isempty(first_step_sb) == true
        plot(xi, yi, ':', 'Color', [1 1 1]);%, ...
            'LineSmoothing', 'on');
    else
        plot(xi, yi, '--', 'Color', [0.5 0.5 0.5]);%, ...
            'LineSmoothing', 'on');
    end
end

if isempty(first_step_sb) == true %draw slipband from the ... center of the grain
    sb_length = distance_segment(xi(1), yi(1), xi(2), yi(2));
    lc = abs(sb_length*cos(2*(pi/4 - (anglesb(sbg) - ...  
        load_orientation))));
    global_dmean(1, 2) = global_dmean(1, 2)+1;
    global_dmean(1, 1) = global_dmean(1, 1)+sb_length;
    if lc>maxlc
        maxlc = lc;
        maxgrainx = xi;
        maxgrainy = yi;
        slip_band_index = sbg;
        slip_band_oriented_angle = anglesb(sbg);
    end
else
    macroangle = get_angle_with_coordinates(xi(1), yi(1), ...  
        xi(2), yi(2));
    ref_angle = get_angle_with_coordinates(first_step_sb(1), ...  
        first_step_sb(2), first_step_sb(3), first_step_sb(4));
    shear_stress_dir = [-3*pi/4-load_orientation ...  
        -pi/4-load_orientation pi/4-load_orientation ...
3*pi/4-load orientation ];
[C, 1] = min(compute_angle_diff(shear_stress_dir, ref_angle));
ref_angle = shear_stress_dir(1);

lc = compute_angle_diff(ref_angle, macroangle);
if lc>=pi
    lc = 2*pi - lc;
end
if lc<= minlc
    minlc = lc;
    maxgrainx = xi;
    maxgrainy = yi;
    slip_band_index = sbg;
    slip_band_oriented_angle = macroangle;
end
end
end

function result = compute_angle_diff(angle1, angle2)
result = abs(sin(angle1) - sin(angle2))+abs(cos(angle1) - cos(angle2));
end

function angle = get_angle_with_coordinates(x1, y1, x2, y2)
    angle = atan((y2 - y1)/(x2 - x1));
% as atan give result between pi/2 and - pi/2, update here:
    if y1<y2 && x1>x2
        angle = angle+pi;
    elseif y1>= y2 && x1>x2
        angle = angle - pi;
    end
end

function [next_grain_id, sb_id, pointx, pointy, sb_length, ...
sb_angle, will_pro] = next_grain_slipband(current_grain, ...
    current_pointx, current_pointy)
will_pro = true;
sb_id = 0;
pointx = 0;
pointy = 0;
sb_length = 0;
sb_angle = 0;

next_grain_id = find_next_grain( V, C, current_grain, ...
    current_pointx, current_pointy );

if next_grain_id ~= 0
    [pointx, pointy, sb_id, sb_angle] = find_slip_band( V, C, ...
        next_grain_id, sb(next_grain_id, :), current_pointx, ...
        current_pointy, ...
        [PL_Structure(1*sign(step)).init_point(1), ...
        PL_Structure(1*sign(step)).init_point(2), ...
        PL_Structure(1*sign(step)).end_point(1), ...
        PL_Structure(1*sign(step)).end_point(2)], ...
        print_slipbands_during_propagation );
else
    will_pro = false;
end

if numel(pointx)>1 && numel(pointy)>1 && will_pro == true
    sb_length = distance_segment(pointx(1), pointy(1), ...
        pointx(2), pointy(2));
else
    will_pro = false;
end

end

[next_grain, next_sb_id, next_points_x, next_points_y, next_sb_length, ...
    next_sb_angle, lado_OK] = next_grain_slipband(...
    PL_Structure(step).grain_id, PL_Structure(step).end_point(1), ...
    PL_Structure(step).end_point(2) );
if next_grain_id == 0 & & numel(next_points_x)>1 & & numel(next_points_y)>1
    PL_Structure(step+sign(step)) = struct('grain_id', next_grain, ...
        'slipband_id', next_sb_id, 'init_point', [next_points_x(1) ...
        next_points_y(1)], 'end_point', [next_points_x(2) ...
        next_points_y(2)], 'length', next_sb_length, 'total_length', ...
        PL_Structure(step).total_length+next_sb_length, 'angle', ...
        next_sb_angle, 'is_border', next_grain == 0);
else
    PL_Structure(step) = struct('grain_id', ...
PL_Structure(step).angle, 'is_border', true);
end
end
end
C.3 File classTM.m

```matlab
classdef classTM
    properties (SetAccess = private, GetAccess = public)
        applied_stress;
        load_direction;
        materialDist = [100];
        sigFL = [368/2];
        surfRoughnessVariance = normrnd(1.0,0.1);
        sigComp = [284];
        nu = [0.29];
        G = [79*10^3];
        propertiesArray = ones(350,8)*(-1);
    end

    methods ( Access = private )
        function [self,res] = getValueByDist(self,i,propertyIndex)
            current_index=1+(1-sign(i))/2;
            if ...
                self.propertiesArray(abs(i),(propertyIndex-1)*2+current_index)==-1
                    if propertyIndex==1
                        matrixValue=self.sigFL;
                    elseif propertyIndex==2
                        matrixValue=self.sigComp;
                    elseif propertyIndex==3
                        matrixValue=self.nu;
                    elseif propertyIndex==4
                        matrixValue=self.G;
                    end
                    %for the first time, compute and store the value
                    if propertyIndex==1
                        matrixValue=self.sigFL;
                    elseif propertyIndex==2
                        matrixValue=self.sigComp;
                    elseif propertyIndex==3
                        matrixValue=self.nu;
                    elseif propertyIndex==4
                        matrixValue=self.G;
                    end
                    %half the crack for both side
                    grainNumber=ceil(1+3*(abs(i)-1)/2);
                    a = randi(100,1,grainNumber); % Martensite: medium carbon steel 0.4C, 142x2
                    c = cell2mat(arrayfun(@(a,matrixValue) ...
                        matrixValue+zeros(1,a), a, matrixValue, 'unif', 0));
                    self.propertiesArray(abs(i),(propertyIndex-1)*2 + ...
                        current_index) = mean(c(:));
                    if (abs(i)==1)%for the first grain, the values are the ...
```
function [self,res] = sigFL(self,i)
    %disp(['sigFL',num2str(i)]);
    [self,res]=self.getValueByDist(i,1);
end

function [self,res] = sigComp(self,i)
    %disp(['sigComp',num2str(i)]);
    [self,res]=self.getValueByDist(i,2);
end

function [self,res] = nu(self,i)
    %disp(['nu',num2str(i)]);
    [self,res]=self.getValueByDist(i,3);
end

function [self,res] = G(self,i)
    %disp(['G',num2str(i)]);
    [self,res]=self.getValueByDist(i,4);
end

function [self,res] = f(self,i)
    [self,sigFLTemp] = self.sigFL(i);
    [self,sigCompTemp] = self.sigComp(i);
    if self.applied_stress > sigFLTemp
        res = 8.3044*10^(-20)*(self.applied_stress)^6.8132;
    else
        res=0;
    end
end

function res = applied_stress(self,i)
    res = self.applied_stress;
end

function [self,res] = phi(self,a,n,i)
    [self,nuTemp] = self.nu(i);
    [self,GTemp] = self.G(i);
    res = ...
\[ 2*(1-\nu_{\text{Temp}})*\sqrt{(1-n^2)}*\text{self.applied stress}(i)*a/(G_{\text{Temp}}*n); \]

    function res = f_1(self,b)
    if(abs(b)<10)
        res=(10-abs(b))/9;
    else
        res=0;
    end
    end

    function res = f_2(self,b)
    if(abs(b)<10)
        res=(abs(b)-1)/9;
    else
        res=1;
    end
    end

    function [self,res] = miml(self,i)
    function res = mi_(t)
        direction_vectors = [
        [1 0 -1],%(-1 1 1)
        [0 -1 1],
        [-1 1 0],
        [0 -1 -1],%(-1 1 1)
        [-1 0 1],
        [1 1 0],
        [1 0 1],%(-1 1 1)
        [-1 0 -1],
        [0 -1 1],%(-1 1 1)
        [-1 1 0],
        [1 1 0]
        ];

        normal_direction_vectors = [
        [1 1 1],%(-1 1 1)
        [1 1 1],
        [1 1 1],
        [1 1 1],%(-1 1 1)
        [1 1 1],
        [1 -1 1],%(-1 1 1)
        [1 -1 1],
        [1 -1 1],
        [1 -1 1],
        [1 -1 1],
        [1 -1 1],
        [1 -1 1]
        ];
    end

APPENDIX C MODEL
function res = normalize(vector)
    length_vec = sqrt(vector(1)^2 + vector(2)^2 + ...
                    vector(3)^2);
    res = [vector(1)/length_vec vector(2)/length_vec ...
                    vector(3)/length_vec];
end

function res = scalar_product(vector1,vector2)
    res = vector1(1)*vector2(1) + ...
         vector1(2)*vector2(2) + vector1(3)*vector2(3);
end

angle=0;
grain_number=ceil(l+3*(abs(t)-1)/2);

if abs(i) > 1
    for a = 1:grain_number
        %random load direction
        load_direction_vec = [random('unif',0.0,2.0)-1 ...
                               random('unif',0.0,2.0)-1 ...
                               random('unif',0.0,2.0)-1];
        orient_matrix=zeros(length(direction_vectors(:,1)),1);
        for k = 1:length(direction_vectors(:,1))
            load_v=normalize(load_direction_vec);
            norm_v=normalize(normal_direction_vectors(k,:));
            dire_v=normalize(direction_vectors(k,:));
            value=abs( 1/( scalar_product( load_v, ...
                              norm_v )*scalar_product( load_v, ...
                              dire_v ) ) );
            orient_matrix(k)=value;
        end
        [m,mi]=sort(orient_matrix);
        lowest5index = mi(1:1);
        angle=angle+mean(orient_matrix(lowest5index));
    end
    res = angle/grain_number;
else
    res = 2.3;
end

end

res = 1 + 2*(mi_(abs(i)) - 2.3) + 2.07*( 2/pi*atan(0.522*{ ...
         abs(i) - 1 })^2 )^-1.86;

end

%constant mim1
function res = mim1_(self,i)
res = 1+2.07*(2/pi*atan(0.522*(abs(i)-1)*2))^-1.86;
end

function [self,res] = sigLi(self,mim1,dmean,c,i)
    [self,sigFLTemp] = self.sigFL(i);
    res = sigFLTemp*mim1*sqrt(dmean/(2*c)) * ... 
         self.surfRoughnessVariance;
end

function [self,res] = nc(self,sigLi,i)
    [self,sigCompTemp] = self.sigComp(i);
    res = cos(pi/2*((self.applied_stress(i)-sigLi)/sigCompTemp));
end

function [self,res] = ns(self,cml,c,nc)
    res = cml/c*nc;
end

function [self,res] = dN(self,ns,nc,i)
    [self,nuTemp] = self.nu(i);
    [self,GTemp] = self.G(i);
    [self,fTemp] = self.f(i);
    if fTemp==0
        res = 0;
    else
        res = ( GTemp/(fTemp*(1 - ... 
                        nuTemp)*2*self.applied_stress(i)) )*(... 
                        asin(nc) - ... 
                        asin(ns));
    end
end

methods ( Access = public )

%constructor
function self = classTM(app_stress,load_direction)
    self.applied_stress = app_stress;
    self.load_direction = load_direction;
end

%i is the current plastic zone side & j the opposite
function [self,res] = ...
    crack_position_by_dN(self,dn,ns,c,i,j, Crack_Structure, PL_Structure)

%the plastic zone is in the entire grain for the first step
if j==0
    j=-sign(i);
end

[self,GTemp] = self.G(i);
[self,nuTemp] = self.nu(i);
[self,fTemp] = self.f(i);

res = abs(c*sin(...
    dN/(GTemp/(fTemp*(1-nuTemp)*2*self.applied_stress(i)))...+
    asin(ns)));
end

% i is the current plastic zone side & j the opposite
function [self,res] = ...
    dadN(self,a,n,i,j,crack_Structure,PL_Structure)

% the plastic zone is in the entire grain for the first step
if j==0
    j=-sign(i);
end

[self,phiTemp] = self.phi(a,n,i);
[self,fTemp] = self.f(i);

res = fTemp*phiTemp;
end

function [dN,nc] = compute_long_crack(self,i,c,dmean,ns)
mim1 = self.mim1(i);
[self,sigFLTemp] = self.sigFL(i);
sigLi = sigFLTemp*mim1*sqrt(dmean/(2*c)) * ...
    self.surfRoughnessVariance;
[self,sigCompTemp] = self.sigComp(i);
nc = cos(pi/2*((self.applied_stress(i)-sigLi)/sigCompTemp));
[self,GTemp] = self.G(i);
[self,nuTemp] = self.nu(i);
[self,fTemp] = self.f(i);

dN = ( GTemp/( fTemp*( 1 - nuTemp ...)
    + 2*self.applied_stress(i) ) )*( asin(nc) - asin(ns) );
end

function matrix = statistics( self, i, ns, mim1_c, length_c, ...
    roughness_c, mim1_v, length_v, roughness_v )
matrix=zeros(4,1);

function value = dN(grain,m,l,r)
    sigLi = sigFLTemp*m*1/r;
    [self,sigCompTemp] = self.sigComp(grain);
    nc = cos(pi/2*(( self.applied_stress(grain) - sigLi ...)
        /sigCompTemp ));
    [self,GTemp] = self.G(grain);
    [self,nuTemp] = self.nu(grain);
    [self,fTemp] = self.f(grain);

    value = (GTemp/( fTemp*(1 - ...
nuTemp)*2* self.applied_stress(grain) ) ...

) + (asin(nc,) - asin(ns));

end

without disp=dN(i,mim1c,lengthc,roughnessc);

matrix(1,1)=dN(i,mim1v,lengthc,roughnessc)-without disp;

matrix(2,1)=dN(i,mim1c,lengthv,roughnessc)-without disp;

matrix(3,1)=dN(i,mim1c,lengthc,roughnessv)-without disp;

matrix(4,1)=0;

end

%does the crack propagate?
% i is the current plastic zone side & j the opposite
function [self,Crack Structure,will pro,disp matrix] = ...

is propagateing(self,i,j,will pro,Crack Structure,PL Structure)

%the plastic zone is in the entire grain for the first step
if j==0

j=-sign(i);

end

dc = PL Structure(i).length;

if abs(i)==1

ddm = PL Structure(1).length+PL Structure(-1).length;

ddm = PL Structure(i).length*2;
else

ddm = dc;
end

%global global_dmean;
%global_dmean(1,1) OR ...
(Crack Structure(i-1*sign(i)).dmean*(abs(i)-1)+ddm)/abs(i)

current step = struct('c', Crack Structure(i-...

1*sign(i)).c+dc,'dmean', (Crack Structure(i-...

1*sign(i)).dmean*(abs(i)-1)+ddm)/abs(i), 'nc', 0, ...

'ac', 0, 'ns', 0, 'dN', 0, 'N', 0, 'is blocked', false);

[self,mim1] = self.mim1(i);

[self,sigLi] = ...

self.sigLi(mim1,current step.dmean,current step.c,i);

[self,current step.nc] = self.nc(sigLi,i);

current step.ac = current step.c+current step.nc;

[self,current step.ns] = self.ns( Crack Structure( i - ......

1*sign(i)).c,current step.c,Crack Structure( i - ......

1*sign(i)).nc );

[self,current step.dN] = ...

self.dN(current step.ns,current step.nc,i);

current step.N = ...
Crack_Structure(i-1*sign(i)).N+current_step.dN;

current_index=1+(1-sign(i))/2;
will_pro(1,current_index) = true;
[self,sigFLTemp]=self.sigFL(i);

%statistics
disp_matrix=zeros(4,1);
disp_matrix=self.statistics(i, current_step.ns, ... 
    self.mim1(i), sqrt(1/(2*abs(i)-1)), l, mim1, ... 
    sqrt(current_step.d_mean/(2*current_step.c)), ... 
    self.surfRoughnessVariance); 

% stop the crack if ... OR if dN==0 or dN<0
if self.applied_stress(i)<sigLi || ... 
    self.applied_stress<sigFLTemp || current_step.dN<0 
    will_pro(1,current_index) = false;
end

current_step.is_blocked = ~will_pro(1,current_index);
Crack_Structure(i) = current_step;
end
C.4 File classFP.m

% classFP.m class
% this is the class for Ferrite-Pearlite

classdef classFP

properties (SetAccess = private, GetAccess = public)
    applied_stress;
    load_direction;
    materialDist = [47 53];%Ferrite & Pearlite
    sigFL = [260/2 385/2];%Ferrite & Pearlite
    surfRoughnessVariance = normrnd(1.0,0.1);
    sigComp = [140 268];%Ferrite: 70*2 & Pearlite: 134*2
    nu = [0.28 0.28];%Ferrite & Pearlite
    G = [82*10^3 82*10^3];%Ferrite & Pearlite
    propertiesArray = ones(350,10)*(-1);
    AVERAGE = 100;
end

methods ( Access = private )

function [self,res] = getValueByDist(self,i,propertyIndex)
    current_index=1+(1-sign(i))/2;
    if ...
        self.propertiesArray(abs(i),(propertyIndex-1)*2+current_index)==-1
            %for the first time, compute and store the value
            if propertyIndex==1
                matrixValue=self.sigFL;
            elseif propertyIndex==2
                matrixValue=self.sigComp;
            elseif propertyIndex==3
                matrixValue=self.nu;
            elseif propertyIndex==4
                matrixValue=self.G;
            elseif propertyIndex==5
                [self,sigFLTemp] = self.sigFL(i);
                if self.applied_stress>sigFLTemp
                    matrixValue=[ ...
                        4.8938*10^-16*(self.applied_stress)^5.4884 ...
                        1.9279*10^-20*(self.applied_stress)^7.0320 ... ];
                else
                    matrixValue=[0 0];
                end
            end
end
end
end
% half the crack for both side
grainNumber=ceil(1+3*(abs(i)-1)/2);
valueToStore=-1;

if abs(i)==1
    valueToStore = matrixValue(1);
elseif abs(i)<self.AVERAGE
    r_ = randi(100,1,grainNumber);
    a = self.materialDist;
    c_ = cell2mat(arrayfun(@(a,matrixValue) ...
        matrixValue+zeros(1,a), a, matrixValue, ...
        'unif', 0));
    valueToStore = mean(c_(r_));
else % the material properties are the average values.
    valueToStore = ( ... matrixValue(1)*self.materialDist(1) + ... ...
        matrixValue(2)*self.materialDist(2) )/100;
end

self.propertiesArray( abs(i), ( propertyIndex - 1 )*2 ... + current_index ) = valueToStore;
if (abs(i)==1)% for the first grain, the values are the same
    self.propertiesArray( abs(i), (propertyIndex-1)*2 ... + 1 + (1-sign(-i))/2 ) = valueToStore;
end

res = self.propertiesArray( abs(i), ... (propertyIndex-1)*2 + current_index );
else %get the value if it was already computed
    res = self.propertiesArray( abs(i), ... (propertyIndex-1)*2 + current_index );
end
end

function [self,res] = sigFL(self,i)
    %disp(['sigFL',num2str(i)]);
    [self,res]=self.getValueByDist(i,1);
end

function [self,res] = sigComp(self,i)
    %disp(['sigComp',num2str(i)]);
    [self,res]=self.getValueByDist(i,2);
end

function [self,res] = nu(self,i)
    %disp(['nu',num2str(i)]);
    [self,res]=self.getValueByDist(i,3);
function [self,res] = G(self,i)
    %disp(['G',num2str(i)]);
    [self,res]=self.getValueByDist(i,4);
end

function [self,res] = f(self,i)
    % [self,sigFLTemp] = self.sigFL(i);
    [self,res]=self.getValueByDist(i,5);
end

function res = appliedstress(self,i)
    res = self.appliedstress;
end

function [self,res] = phi(self,a,n,i)
    [self,nuTemp] = self.nu(i);
    [self,GTemp] = self.G(i);
    res = ...
        2*(1-nuTemp)*sqrt(1-nˆ2)*self.appliedstress(i)*a/(GTemp*n);
end

function [self,res] = mim1(self,i)
    function res = mim(t)
        direction_vectors = [
            [1 0 -1],% (1 1 1)
            [0 -1 1],
            [-1 1 0],
            [0 -1 -1],% (1 -1 1)
            [-1 0 1],
            [1 1 0],
            [1 1 0],% (-1 1 1)
            [-1 0 -1],
            [0 -1 1],% (-1 -1 1)
            [-1 1 0],
            [1 0 1]
        ];
        normal_direction_vectors = [
            [1 1 1],% (1 1 1)
            [1 1 1],
            [1 1 1],
            [1 -1 1],% (1 -1 1)
            [1 -1 1],
            [1 -1 1],% (-1 1 1)
            [-1 1 1],
            [-1 1 1],
            [-1 1 1]
        ];
    end
function res = normalize(vector)
    length_vec = sqrt(vector(1)^2 + vector(2)^2 + vector(3)^2);
    res = [vector(1)/length_vec vector(2)/length_vec ... vector(3)/length_vec];
end

function res = scalar_product(vector1, vector2)
    res = vector1(1)*vector2(1) + vector1(2)*vector2(2) ... + vector1(3)*vector2(3);
end

angle = 0;
grain_number = ceil(1 + 3*(abs(t) - 1)/2);

if abs(i) > 1
    for a = 1:grain_number
        % random load direction
        load_direction_vec = [random('unif', 0.0, 2.0)-1 ... random('unif', 0.0, 2.0)-1 ... random('unif', 0.0, 2.0)-1];

        orient_matrix = zeros(length(direction_vectors(:,1)), 1);
        for k = 1:length(direction_vectors(:,1))
            load_v = normalize(load_direction_vec);
            norm_v = normalize(normalize(direction_vectors(k,:)));
            dire_v = normalize(direction_vectors(k,:));
            value = abs(1/scalar_product(load_v, ... norm_v)*scalar_product(load_v, ... dire_v));
            orient_matrix(k) = value;
        end

        [m, mi] = sort(orient_matrix);
        lowest5index = mi(1:5);
        angle = angle + mean(orient_matrix(lowest5index));
    end
    res = angle/grain_number;
else
    res = 2.3;
end
end

res = 1 + 2*( mi(abs(i)) - 2.3 ) + 2.07* ... 2/pi*atan(0.522*( abs(i) - 1 )^2)^1.86;
function res = mim1(self,i)
    res = 1+2.07*(2/pi*atan( 0.522*(abs(i)-1)*2))ˆ1.86;
end

function [self,res] = sigLi(self,mim1,dmean,c,i)
    [self,sigFLTemp] = self.sigFL(i);
    res = sigFLTemp*mim1*sqrt(dmean/(2*c)) + ... 
         self.surfRoughnessVariance;
end

function [self,res] = sigLi(self,mim1,dmean,c,i)
    [self,sigCompTemp] = self.sigComp(i);
    res = cos(pi/2*((self.applied_stress(i)-sigLi)/sigCompTemp));
end

function [self,res] = ns(self,cml,c,nc)
    res = cml/c*nc;
end

function [self,res] = dN(self,ns,nc,i)
    [self,nuTemp] = self.nu(i);
    [self,GTemp] = self.G(i);
    [self,fTemp] = self.f(i);
    if fTemp== 0
        res = 0;
    else
        res = ( GTemp/(fTemp*( 1 - nuTemp ... 
                        +2*self.applied_stress(i)) )*( asin(nc) - ... 
                        asin(ns) ));
    end
end

methods ( Access = public )

%constructor
function self = classFP(app_stress,load_direction)
    self.applied_stress = app_stress;
    self.load_direction = load_direction;
end

%i is the current plastic zone side & j the opposite
function [self,res] = ...
    crack.position_by_dN(self,dN,ns,c,i,j,Crack.Structure,PL_Structure)

%the plastic zone is in the entire grain for the first step
if j==0
    j=-sign(i);
end
[self,GTemp] = self.G(i);
[self,nuTemp] = self.nu(i);
[self,fTemp] = self.f(i);

res = abs(c*sin( ... 
dN/(GTemp/(fTemp*(1-nuTemp)+2*self.applied_stress(i)) ... 
+ asin(ns))));
end

% i is the current plastic zone side & j the opposite
function [self,res] = ... 
dadN(self,a,n,i,j,Crack_Structure,PL_Structure)
%the plastic zone is in the entire grain for the first step
if j==0
  j=-sign(i);
end

[self,phiTemp] = self.phi(a,n,i);
[self,fTemp] = self.f(i);
res = fTemp*phiTemp;
end

function [dN,nc] = compute_long_crack(self,i,c,dmean,ns)
mim1 = self.mim1(i);
% here i=self.AVERAGE to have average values
[self,sigFLTemp] = self.sigFL(self.AVERAGE);
sigLi = sigFLTemp*mim1*sqrt(Crack_Structure(self.AVERAGE)/(2*c)) * ... 
  self.surfRoughnessVariance;
[self,sigCompTemp] = self.sigComp(self.AVERAGE);
nc = cos(pi/2*((self.applied_stress(i)-sigLi)/sigCompTemp));
[self,GTemp] = self.G(self.AVERAGE);
[self,nuTemp] = self.nu(self.AVERAGE);
[self,fTemp] = self.f(self.AVERAGE);
dN = (GTemp/(fTemp*(1- ... 
nuTemp+2*self.applied_stress(i)))*( asin(nc) - ... 
  asin(ns)))
end

function matrix = statistics(self, i, ns, mim1,c, length_c, ... 
  roughness_c, mim1_v, length_v, roughness_v)
matrix=zeros(4,1);

function value = dN(grain,m,l,r)
[self,sigFLTemp] = self.sigFL(grain);
sigLi = sigFLTemp*m[l/r];
[self,sigCompTemp] = self.sigComp(grain);
nc = cos( pi/2*((self.applied_stress(grain) - ... 
  sigLi)/sigCompTemp) );
[self,GTemp] = self.G(grain);
[self,nuTemp] = self.nu(grain);
[self,fTemp] = self.f(grain);

value = (GTemp/(fTemp*( 1 - nuTemp ... )^2)*self.applied_stress(grain))*/( asin(nc) - ... 
        asin(ns) );

end

without disp=dN(self.AVERAGE,mim1c,length_c,roughness_c);
matrix(1,1)=dN(self.AVERAGE,mim1v,length_c,roughness_c)-without disp;
matrix(2,1)=dN(self.AVERAGE,mim1c,length_v,roughness_c)-without disp;
matrix(3,1)=dN(self.AVERAGE,mim1c,length_c,roughness_v)-without disp;
matrix(4,1)=dN(i,mim1c,length_v,roughness_c)-without disp;

end

%does the crack propagate?
% i is the current plastic zone side & j the opposite
function [self,Crack_Structure,will pro,disp_matrix] = ...
    is propagating(self,i,j,will pro,Crack Structure,PL Structure)

%the plastic zone is in the entire grain for the first step
if j==0
    j=-sign(i);
end

dc = PL Structure(i).length;
if abs(i)==1
    ddm = PL Structure(1).length+PL Structure(-1).length;
else
    ddm = dc;
end

%global global dmean;
%global global dmean
(Crack Structure(i-1*sign(i)).d_mean*(abs(i)-1)+ddm)/abs(i)
current_step = struct('c', Crack Structure(i - ... 
    1*sign(i)).c+dc,'d_mean', (Crack Structure(i - ... 
    1*sign(i)).d_mean*(abs(i) - 1)+ddm)/abs(i), 'nc', 0, ...
    'ac', 0, 'ns', 0, 'dN', 0, 'N', 0, 'is blocked', false);

[self,mim1] = self.mim1(i);
[self,sigLi] = ...
        self.sigLi(mim1,current_step.d_mean,current_step.c,i));
[self,current_step.nc] = self.nc(sigLi,i);

current_step.ac = current_step.c*current_step.nc;
[self,current_step.ns] = self.ns( Crack Structure( i - ... 
    1*sign(i) ).c,current_step.c, Crack Structure( i - ...
1*sign(i) ).nc );
self.derivative = self.derivative;  

[ self, current_step.dN ] = ...
self.derivative = self.derivative;  

current_step.dN = ...
Crack_Structure(i-1*sign(i)).N+current_step.dN;

current_index=l+(1-sign(i))/2;
will_pro(1,current_index) = true;
[ self, sigFLTemp ] = self.sigFL(i);

% statistics
disp_matrix=zeros(4,1);
disp_matrix = self.statistics(i, current_step.ns, ...  
self.mim1, 1, mim1, ...  
sqrt(current_step.d_mean/(2*current_step.c)), ...  
surfRoughnessVariance_ );

% stop the crack if ... OR if dN==0 or dN<0
if self.applied_stress(i)<sigLi || ...  
self.applied_stress<sigFLTemp || current_step.dN<=0
will_pro(1,current_index) = false;
end

current_step.is_blocked = ~will_pro(1,current_index);
Crack_Structure(i) = current_step;

end

end

end
Appendix C presents Scanning Electron Microscopy (SEM) images used for the 3D reconstruction of the surface roughness profile with photometric stereo technique.

**Figure D.1** – 3D surface reconstruction using photometric stereo technique: (a)-(d) source SEM images.
Figure D.1 – 3D surface reconstruction using photometric stereo technique: (e) 3D reconstructed surface.
S-N-P fatigue curves using maximum likelihood
Method for fatigue resistance curves with application to straight and welded rebars

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Introduction

When checking the fatigue life of steel bridges with concrete deck slabs, both steel details and embedded reinforcing steel bars (rebars) must be considered. In this paper a method for estimation of fatigue resistance curves is presented with application to straight and welded rebars. Rebar fatigue resistance is traditionally presented in the form of $S-N-P$ curves which relate the applied stress range, $S$, to the $p$-quantile of fatigue life, $N$. These $S-N-P$ curves are obtained from rebar fatigue tests at constant stress amplitudes. Test results for hot rolled (HR), cold worked (CW) and quenched and self-tempered (QST) rebars can be found in [1-7]. Both HR and CW rebars show a ductile cross section consisting of pearlite-ferrite microstructure and low or medium Carbon content [8, 9]; however, HR and CW rebars have been mainly replaced by QST rebars since the 1970’s. QST rebars are produced from a specific thermal treatment called Thermex or Tempcore [10, 11] which results in a different microstructure at the surface and in the core. Typically the surface is a hard martensite, whereas the core consists of pearlite-ferrite. Fatigue datasets for different rebar connections such as lapping, coupling or welding as well as for corroded rebars can be found from tests performed in the 1970’s.

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In the EN standards, characteristic S-N curves are created by fitting a linear regression to the experimental failure data points and translating the linear regression mean curve to the p-lower hyperbolic prediction bound (typically $p=0.05$), at 1 million cycles [12]. This approach has several limitations: 1) run-out test results are neglected; 2) a constant amplitude fatigue limit (CAFL) (stress below which tested bars experience no fatigue damage) is arbitrarily chosen to begin at 1 million cycles for straight rebars and at 10 million cycles for welded rebars [13]; and 3) S-N curves are based on fatigue data scatter in the finite-life region ($N$ less than 1 million cycles) resulting in less accuracy in the high cycle fatigue (HCF) region ($N$ over 1 million cycles). The statistical method recommended by the EN standards is currently used in the standards for concrete structures [14]. Analysing the fatigue data using more statistically robust approaches may overcome some of these issues.

In this paper a Maximum Likelihood (ML) method-based approach is used together with Monte-Carlo Simulations (MCS) to estimate $S$-$N$-$P$ curves of straight and welded rebars. Run-out test results are considered and particular attention is given to the position of the CAFL. The influence of rebar size and rebar type is studied. Comparisons between ML-based $S$-$N$-$P$ curves and EN-based characteristic S-N curves are made. The approach proposed in this study is presented with application to straight and welded rebars but it has generic applicability for estimating fatigue $S$-$N$-$P$ curves of fatigue sensitive details of steel bridges like welded and bolted connections.

E.1 Statistical evaluation of S-N-P curves

This section presents the statistical method for estimation of characteristic S-N curves recommended by the EN standards [12], and presents the formulation for the ML-based approach.

E.1.1 Statistical evaluation of S-N-P curves based on EN background documentation

A linear statistical model is used to define the relationship between the logarithm of the number of cycles to failure, $Y = \ln(N)$, and the logarithm of the nominal stress range, $X = \ln(S)$:

$$Y = \beta_0 + \beta_1 \cdot X + \epsilon(0, \sigma) \quad (E.1)$$

In Eq. 1, $\beta_0$ and $\beta_1$ are respectively the intercept and the slope of the S-N curve in
the log-log plane. It is assumed that the model error $\epsilon$ can be modeled with a normal random variable, with an expected value equal to zero and standard deviation equal to $\sigma$. The model $E(Y) = \beta_0 + \beta_1 \cdot X$, which represents the mean value of $\ln N$ for an assigned stress range, is fitted to the experimental dataset $(y_1, x_1) \ldots (y_n, x_n)$ using the least square method (LSM). Only failure points are considered.

According to the LSM, the unbiased, normally distributed estimators of model parameters are:

$$\hat{\beta}_1 = \frac{\sum_{i=1}^{n} (x_i - \bar{x})(y_i - \bar{y})}{\sum_{i=1}^{n} (x_i - \bar{x})^2} = \frac{S_{xy}}{S_{xx}} \quad \text{(E.2)}$$

$$\hat{\beta}_0 = \bar{y} - \hat{\beta}_1 \bar{x} \quad \text{(E.3)}$$

Since $\hat{\beta}_0$ and $\hat{\beta}_1$ are normally distributed in repeated sampling, it follows that $\hat{N} = \hat{\beta}_0 + \hat{\beta}_1 \cdot X$ is also normally distributed. In order to obtain the characteristic S-N curve, a 100(α)% lower hyperbolic prediction bound can be determined around the mean regression line, using the following expression:

$$\hat{\beta}_0 + \hat{\beta}_1 \cdot x^* + t_{\alpha,n-2} \cdot StD \cdot \sqrt{1 + \frac{1}{n} + \frac{(x^* - \bar{x})^2}{S_{xx}}} \quad \text{(E.4)}$$

In Eq. 4, $StD$ is the sample standard deviation, $t_{\alpha,n-2}$ is the $\alpha$-quantile of the Student’s T distribution with $n-2$ degrees of freedom and $x^*$ is the natural logarithm of the reference stress range. Characteristic S-N curves are determined by translating the mean regression line to the corresponding point of the 5% lower hyperbolic prediction bound, at 1 million cycles. The constant amplitude fatigue limit (CAFL) is arbitrarily chosen to begin at 1 million cycles for straight bars and at 10 million cycles for welded bars [13].

### E.1.2 Statistical evaluation of S-N-P curves based on maximum likelihood approach

As previously mentioned, this EN standard approach is limited because: 1) run-out test results are neglected (loss of information) 2) the CAFL position is arbitrarily chosen; and 3) prediction bounds of linear regression curves are based on fatigue data scatter in the finite-life region resulting in less accuracy in the HCF region. To overcome these limitations, Pascual et al. [15] proposed a 5-parameter random fatigue limit (RFL) model that fit a nonlinear S-N curve having a random CAFL, to a complete fatigue dataset using ML estimation. In [15], characteristic S-N curves
were determined by finding lower \( \alpha \)-confidence bounds of \( p \)-quantile S-N curves (typically \( \alpha=75\% \) and \( p=0.05 \)). This ML model proposed by Pascual is still affected by two limitations: 1) the choice of the \( \alpha \)-confidence level for the lower bound of the \( p \)-quantiles is arbitrary; and 2) RFL-based S-N curves are nonlinear and are not easily comparable to the current standard linear S-N curves from the EN method.

This study proposes a bi-linear random fatigue limit (BLRFL) model that fits a bi-linear median S-N curve to a complete fatigue dataset, using again ML estimation. S-N-P curves are computed using Monte-Carlo Simulations (MCS), whereby the arbitrary choice of the \( \alpha \)-confidence level for the lower bound of the \( p \)-quantiles is not required.

The dependence between fatigue life and stress range is modeled as follows:

\[
Y = \frac{\beta_0 + \beta_1 \cdot X}{H(X - V)} + \epsilon (0, \exp(\sigma_Y)) \quad \text{(E.5)}
\]

Where \( H(\cdot) \) is the unit step function and \( V \) is the natural logarithm of CAFL. \( Y \) and \( V \) are assumed to be normal distributed random variables:

\[
Y = \text{Normal}(\mu_Y, \exp(\sigma_Y)) \quad \text{(E.6)}
\]

\[
V = \text{Normal}(\mu_V, \exp(\sigma_V)) \quad \text{(E.7)}
\]

The location parameter of the \( Y \) distribution is:

\[
\mu_Y = \frac{(\beta_0 + \beta_1 \cdot X)}{H(X - V)} \quad \text{(E.8)}
\]

The conditional probability density function of \( Y|V \) is:

\[
f_{Y|V} = \frac{1}{\sigma_Y} \phi_{Y|V}(x, y, v; \beta_0, \beta_1, \sigma_Y) \quad \text{(E.9)}
\]

The marginal probability density function of \( Y \) is:

\[
f_Y = \int_{-\infty}^{\infty} \frac{1}{\sigma_Y} \phi_{Y|V}(x, y, v; \beta_0, \beta_1, \sigma_Y) \phi_V(v; \mu_V, \sigma_V) \, dv = f_Y(x, y; \theta) \quad \text{(E.10)}
\]

Similarly the marginal cumulative distribution function of \( Y \) is:

\[
F_Y = \int_{-\infty}^{x} \frac{1}{\sigma_Y} \phi_{Y|V}(x, y, v; \beta_0, \beta_1, \sigma_Y) \phi_V(v; \mu_V, \sigma_V) \, dv = F_Y(x, y; \theta) \quad \text{(E.11)}
\]
Where the model parameter vector is indicated as:

$$\begin{align*}
\underline{\theta} &= (\beta_0, \beta_1, \sigma_Y, \mu_V, \sigma_V) \quad (E.12)
\end{align*}$$

The sample likelihood is:

$$L(\theta) = \prod_{i=1}^{N_{\text{fail}}} \left[ f_Y(x_i, y_i; \theta) \right]^{\delta_i} \left[ 1 - F_Y(x_i, y_i; \theta) \right]^{1-\delta_i} \quad (E.13)$$

where $\delta_i = 1$ for the failure points and $\delta_i = 0$ for run-out points. The negative sample log-likelihood is:

$$-\ln(L(\theta)) = - \left( \sum_{i=1}^{N_{\text{fail}}} \ln(f_Y(x_i, y_i; \theta)) + \sum_{i=1}^{N_{\text{runouts}}} \ln(1 - F_Y(x_i, y_i; \theta)) \right) \quad (E.14)$$

The maximum likelihood estimate of the parameters vector $\underline{\theta}$ is the vector that minimizes the negative sample log-likelihood.

$$E(\underline{\theta}) = [E(\beta_0), E(\beta_1), E(\sigma_Y), E(\mu_V), E(\sigma_V)] \quad (E.15)$$

The inverse of the Fisher information matrix is the asymptotic covariance matrix $C$ and gives information on the uncertainty of the stochastic model. Once the vector $E(\underline{\theta})$ and the covariance matrix $C$ have been computed, following MCS approach is used to estimate S-N-P curves:

- A stress range $S$ is selected
- $10^6$ values of $Y$ are sampled using $E(\underline{\theta})$ and $C$ information
- $P_t$ is computed for each value of the sample
- the $p$-quantile of the fatigue life, $N$ that gives $E(P_t)=p$, for the selected stress range
- the process is repeated

### E.2 Results of statistical analysis

Five different experimental datasets were analysed using both the EN- and ML-based approaches. The five different data sets represent:

- HRCW straight rebars with diameter, $d$, smaller than 20 mm [1, 2]
- HRCW straight rebars with $d$ greater than 20 mm [1–3, 7]
- QST straight rebars with $d$ smaller than 20 mm [4][anonymous industrial dataset]
- QST straight rebars with \( d \) greater than 20 mm \([4, 5]\)
- HR butt welded rebars (60-degree single-V weld joint) \([6]\)

Figs. 1 to 3 (b) show the ML-based median S-N curves, the ML-based 5\(^{th}\) quantile S-N curves and EN-based characteristic curves for the five considered datasets. ML-based 5\(^{th}\) quantile nonlinear S-N curves were linearized (dotted lines) for direct comparison with EN-based characteristic S-N curves, using the following approach: 1) a horizontal line at CAFL\(_{5\%}\) is traced; 2) a straight line with slope equal to the slope of the median line and starting at the lower abscissa point of the nonlinear curve is intersected with the CAFL\(_{5\%}\) horizontal line; 3) the intersection point is the knee point. For all considered datasets, the ML-based 5\(^{th}\) quantile S-N curves and EN-based characteristic curves are almost identical while \( N < 10^6 \) cycles.

![Diagram 1](image1)

**Fig. 1.** S-N curves for HRCW straight rebars; (a) diameter \( \leq 20 \text{ mm} \); (b) diameter > 20 mm

Figs. 1 and 2 show that for HRCW and QST straight rebars ML-approach gives considerably lower estimation of the CAFL with respect to the EN-based approach:
ML-approach gives estimates of the knee point between 1.4 and 3.4 million cycles (see Table E.1).

Fig. 3 (a) shows that for HR welded rebars the EN-based approach gives an over conservative estimate of the CAFL of the characteristic curve with respect to the ML-based approach: ML-approach gives estimate of the knee point at 5.5 million cycles (see Table E.1).

For HRCW and QST straight rebars the fatigue resistance increases by decreasing the diameter of the section; for HRCW straight rebars the CAFL of the ML-based 5\(^{th}\) quantile S-N curve decreases from 162 MPa (d\(_{\leq} 20\) mm) to 134 MPa (d\(_{>20}\) mm) while the fatigue resistance of the ML-based 5\(^{th}\) quantile curve at \(10^6\) cycles decreases from 224 MPa (d\(_{\leq} 20\) mm) to 152 MPa (d\(_{>20}\) mm). For QST straight rebars, the CAFL of the ML-based 5\(^{th}\) quantile S-N curve decreases from 214 MPa (d\(_{\leq} 20\) mm) to 188 MPa (d\(_{>20}\) mm) while the fatigue resistance of the ML-based 5\(^{th}\) quantile curve at \(10^6\) cycles decreases from 234 MPa (d\(_{\leq} 20\) mm) to 107 MPa (d\(_{>20}\) mm) (see Table E.1).

![Graph showing S-N curves for QST straight rebars](image)

Fig. 2. S-N curves for QST straight rebars; (a) diameter \(_{\leq} 20\) mm; (b) diameter \(_{>20}\) mm
Table E.1 – Summary of characteristic values of ML-based linearized S-N curves

<table>
<thead>
<tr>
<th>Type of rebars</th>
<th>Slope</th>
<th>$\text{CAFI}_{50%}$</th>
<th>$\text{CAFI}_{5%}$</th>
<th>knee point</th>
<th>$S (N=10^6)_{5%}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>HRCW $d \leq 20$ mm</td>
<td>-5.17</td>
<td>219 MPa</td>
<td>162 MPa (200 MPa)</td>
<td>$3.4 \times 10^6$</td>
<td>224 MPa</td>
</tr>
<tr>
<td>HRCW $d &gt; 20$ mm</td>
<td>-4.72</td>
<td>146 MPa</td>
<td>134 MPa (144 MPa)</td>
<td>$1.4 \times 10^6$</td>
<td>152 MPa</td>
</tr>
<tr>
<td>QST $d \leq 20$ mm</td>
<td>-4.46</td>
<td>279 MPa</td>
<td>214 MPa (240 MPa)</td>
<td>$1.8 \times 10^6$</td>
<td>258 MPa</td>
</tr>
<tr>
<td>QST $d &gt; 20$ mm</td>
<td>-4.39</td>
<td>233 MPa</td>
<td>188 MPa (225 MPa)</td>
<td>$2.3 \times 10^6$</td>
<td>234 MPa</td>
</tr>
<tr>
<td>Welded</td>
<td>-2.71</td>
<td>123 MPa</td>
<td>48 MPa (34 MPa)</td>
<td>$5.5 \times 10^6$</td>
<td>107 MPa</td>
</tr>
</tbody>
</table>

Fig. 3. (a) S-N curves for HR welded rebars (b) ML 5$^{th}$ quant. linearized S-N curves for all experimental datasets

ML-based 5$^{th}$ quantile S-N curves were plotted in Fig. 3 for the five considered datasets: both for $d \leq 20$ mm and for $d > 20$ mm QST straight rebars show higher fatigue resistance with respect to HRCW straight rebars. HR welded rebars have the by far lowest fatigue resistance both in terms of CAFL and of stress range at $10^6$ cycles.
Figs. 2 and 3 (a) show that QST straight rebars and HR welded rebars have a small deviation of \( 5^{\text{th}} \) quantile curve from the median curve in the finite life region and higher deviation in HCF region. Fig. 1 (a) shows that HRCW \((d>20 \text{ mm})\) straight rebars have high deviation both in finite life region and in HCF region; HRCW \((d>20 \text{ mm})\) straight rebars have high deviation in finite life region and a smaller deviation in HCF region.

### E.3 Discussion of statistical analysis results

Comparison of the ML-based S-N curves and the EN-based characteristic curves in the HCF region for HRCW and QST straight rebars, indicates that the arbitrary assumption of having the CAFL at \(10^6\) cycles is unsafe since ML estimates of the S-N curve knee point lie between 1.4 and 3.4 million cycles. On the contrary, comparison of ML-based S-N curves and EN-based characteristic curves in the HCF region for HR welded rebars, indicate that the arbitrary assumption of having the CAFL at \(10^7\) cycles seems is over conservative since the ML estimate of knee point of the S-N curve lies at 5.5 million cycles.

ML-based linearized \(5^{\text{th}}\) quantile S-N curves indicate that fatigue resistance of HRCW and QST straight rebars decreases as the diameter increases. For a given diameter interval, QST straight rebars have higher fatigue resistance with respect to HRCW straight rebars. HR welded rebars have the lowest fatigue resistance within all analysed datasets.

High deviation of the ML-based \(5^{\text{th}}\) quantile curve from the median curve was observed in HCF region for QST straight rebars, HRCW \((d<20 \text{ mm})\) straight rebars and HR welded rebars: this is due to the fact that the experimental datasets are highly dispersed in the HCF region. On the contrary a small standard deviation was observed in HCF region for HRCW straight \((d>20 \text{ mm})\) rebars, which is probably due to the fact that only run-out points exist at lowest stress ranges in the experimental datasets.

In conclusion the findings of this paper suggest that the limitations included in the current EN recommendations for statistical evaluations of characteristic S-N curves lead to incoherent fatigue resistance estimation in the HCF region for all types of analysed rebars. The ML-approach proposed herein constitutes a powerful tool that can be used to re-define the characteristic S-N curves for straight and welded rebars by taking in account both rebar type and size effect. The estimation of the characteristic S-N curves in the HCF region is directly related to experimental
data and the coherence of the estimates can be ameliorated by increasing the significance of the dataset information in the HCF region. Furthermore it has to be noted that the ML-approach is presented with application to straight and welded rebars but it has generic applicability for estimating fatigue S-N-P curves of fatigue sensitive details of steel bridges like welded and bolted connections.

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Bibliography


Appendix F

Curriculum Vitae

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Education

2010–2014 PhD in Structural Engineering, Swiss Federal Institute of Technology in Lausanne – EPFL, Lausanne, Switzerland
- Thesis entitled: Fatigue Behaviour of Steel Reinforcement Bars at Very High Number of Cycles
- Directors: Prof. E. Brühwiler and Prof. A. Nussbaumer

2007–2010 Master of Science in Structural Engineering, Polytechnic School at the University of São Paulo – EPUSP, São Paulo - SP, Brazil
- Director: Prof. Dr. Marcos Aurélio Marques Noronha

2002–2007 Bachelor in Civil Engineering, Federal University of Piauí – UFPI, Teresina - PI, Brazil
- Graduate Research: Structural analysis of plane frames for infrastructure project - comparison between classical methods and Finite Element Method.

Teaching and Professional Experiences

2011–2013 Teaching EPFL, Lausanne, Switzerland
- Description: Teaching assistant for courses in Existing structures, structural safety & reliability and aesthetics of structures at the School of Architecture, Civil and Environmental Engineering (ENAC) of EPFL. Supervision of master and undergraduate projects at EPFL.
**Structural Engineer**  
Celenge Engenharia, São Paulo - SP, Brazil  
Description: Structural calculation of bridges; design of slabs, beams and calculation of forms. Checking of conditions of stability and design of retaining walls.

**Internship as Site Manager**  
Construtora CAXÉ, Teresina - PI, Brazil  
Description: Supervision of building sites such as schools, squares and housing estates. Update schedules of work with the physical and financial progress and reporting of the work week.

**Internship as Structural Engineer**  
Maloca Arquitetura e Estruturas, Teresina - PI, Brazil  
Description: Structural calculation of infrastructure projects such as houses, with design of slabs, beams. Structural calculation of bridges with design of foundations.

**Honors and Awards**

2014  
3rd prize  
ENAC Doctoral Poster, EPFL

2008-2009  
Postgraduate research scholarship  
FAPESP

2005-2006  
Graduate research scholarship  
Institutional Program of Scientific Initiation

2006  
Poster selected for presentation  
XV Seminar of Scientific Initiation, UFPI

**Journal Papers**

2014  

2014  

2014  

2014  
Conference Proceedings

2014 D'Angelo, L.; Rocha, M.; Nussbaumer, A. & Brühwiler, E. S-N-P fatigue curves using maximum likelihood. 7th Conference on Steel and Composite Structures, Napoli, Italy (accepted).


2009 Nunes, M. R. P. P.; Noronha, M. A. M. Um novo algoritmo para modelagem de Mecânica da Fratura usando o Método dos Elementos de Contorno. XXX Iberian Latin American Congress on Computational Methods in Engineering (CILAMCE), Búzios, Brazil.

Languages

Portuguese, mother tongue
English, fluent
French, intermediate

Computer Skills

Document Preparation System: \LaTeX, Microsoft Office (Word, Excel, PowerPoint)
Finite Element Software: ABAQUS
Computer Programming: Java, Visual Basic & VBA, Delphi, Netbeans, Eclipse, UML
Numerical Computing: Matlab
Computer-Aided Design: AutoCAD
Vector Graphics Editor Application: Inkscape