

# The Effects of Quenching and Tempering Treatment on the Hardness and Microstructures of a Cold Work Steel

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*Abstract. The X153CrMoV12 ledeburitic chromium steel characteristically has high abrasive wear resistance, due to their high carbon and high chromium contents with a large volume of carbides in the microstructure. This steel quality has high compression strength, excellent deep hardenability and toughness properties, dimensional stability during heat treatment, high resistance to softening at elevated temperatures. The higher hardness of cryogenic treated samples in comparison with conventional quenched samples mean lower quantity of retained austenite as at samples quenched to room temperature and tempered in similar condition. In the microstructure of samples were observed that the primary carbide did not dissolve at 1070°C and their net structure have not been changed during to heat treatment. During to tempering at high temperature the primary carbides have become more and more rounded. After low tempering temperature in martensite were observed some small rounded carbides also, increasing the tempering temperature the quantity of finely dispersed carbides increased, which result higher hardness. The important issues in heat treatment of this steels are the reduction or elimination of retained austenite due to cryogenic treatment.*

## Introduction

Tool steels are divided into different category as: cold work, hot work, plastic mould, high speed tool steels [1, 2]. In many industrial applications tool steels are subjected to extremely high and variable loads. Cold work tool steels are used for fabricating stamping or forming dies, punches, forming rolls, knives, slitters, shear blades or any other component for shaping a material into a part or component adapted to a definite use. The mechanical properties of these steels can be improved by modifying of microstructure. The microstructure can be modified by using different heat treatments technologies.

Tool steel properties and its wear resistance can be enhanced by optimizing heat treatment parameters [3]. This type of steels is generally used in a quenched and tempered state and, therefore, the precipitation of secondary carbides and the evolution of the matrix in the steel during tempering determine the properties of the steel. The tempering process at unalloyed and low alloyed steels usually leads to a decrease in strength due to the precipitation of carbides from carbon that is originally in solid solution in the martensite.

Increasing the carbon content and alloy content of the steel such as by increasing the austenitising temperature and holding time, will produce a harder martensite, but simultaneously lower the  $M_s$  and  $M_f$  temperatures. After Hollomon and Jaffe [4] this temperature can be calculated

$$M_s = 550 - 350C\% - 40Mn\% - 35V\% - 20Cr\% - 17Ni\% - 10Cu\% - 10Mo\% - 8W\% + 15Co\% + 30Al\% \quad (1).$$

Hougardy proposed a correction to this formula [5]:

$$M_s = 0.495M_{sjh} + 0.00095M_{sjh}^2 + 40 \quad (2)$$

$M_s$  - Corrected martensite start temperature [°C]

$M_{sjh}$  - Martensite temperature start according to Jaffe & Hollomon [°C]

*Alloy amount* - [weight %]

After Hougardy the volume fraction of martensite can be calculated by

$$V_M = 1 - \exp[-k(M_s - T)^q] \quad (3)$$

$$k = 0.36 \times 10^{-3} + 0.10 \times 10^{-4} M_s - 0.34 \times 10^{-6} M_s^2 + 0.32 \times 10^{-8} M_s^3 - 0.52 \times 10^{-11} M_s^4 \quad (4)$$

$$q = 2.088 - 0.76 \times 10^{-2} M_s + 0.16 \times 10^{-4} M_s^2 - 0.90 \times 10^{-8} M_s^3 \quad (5)$$

Notation:

$V_M$  - Volume fraction of martensite

$T$  - temperature [°C]

$M_s$  - Temperature at which 1% martensite forms [°C]

The presence of high carbon and high alloy content in tool steels lower their characteristic temperatures of martensite start and martensite finish [6]. Therefore, conventional hardening treatment of these steels fails to convert considerable amount of austenite into martensite often leading to unacceptable level of retained austenite in the as quenched structure of these steels. The retained austenite is soft and thus adversely affects the desirable properties such as hardness and wear resistance [7]. Moreover, the retained austenite is unstable and transforms into martensite at the service conditions of tool steels. The freshly formed martensite being untampered is very brittle and hence undesirable. Furthermore, transformation of austenite to martensite is associated with approximately 4% volume expansion [8], which leads to dimensional changes and distortion of the components, even failure in extreme cases [9].

Therefore, one of the major challenges in the heat treatment of tool steels is to minimize the amount of retained austenite or eliminate it.

Retained austenite content in tool steels can be reduced substantially by cryogenic treatment [10]. Multiple tempering and cryogenic treatments improves the strength and hardness of tool steels.



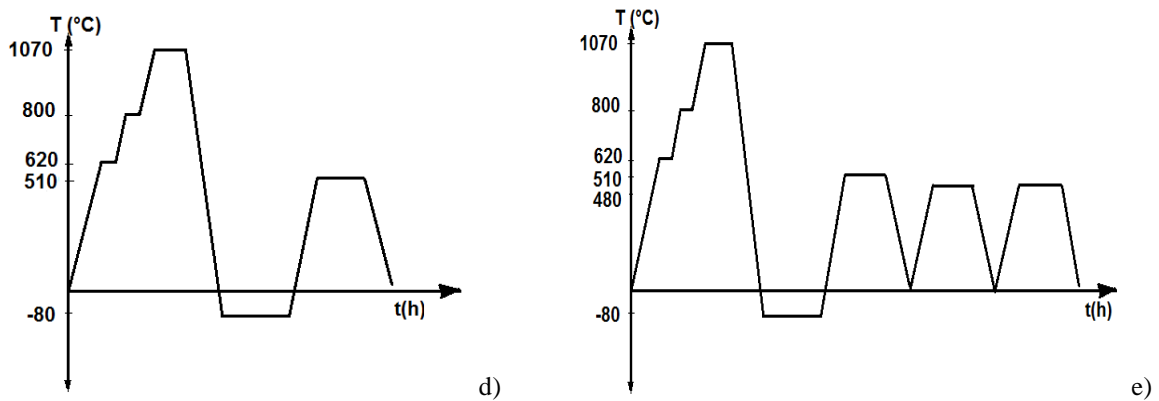


Figure 1. Heat treatments diagrams of specimens a) nr.1 b) nr. 2 c) nr.3 d) nr.4 e) nr.5

The microhardness testing were executed with Vickers testing machine type Buhler 1105. The microstructures of samples were studied by Olympus PMG3 microscope after conventional metallographic preparation. For etching were used nital 2%.

The microstructure of samples in delivered condition consist primary carbides on borders of the primary grains in the matrix with spheroid structure as it is showed in *Figure 2*. The hardness of samples in this conditions were 240HV<sub>1</sub>.

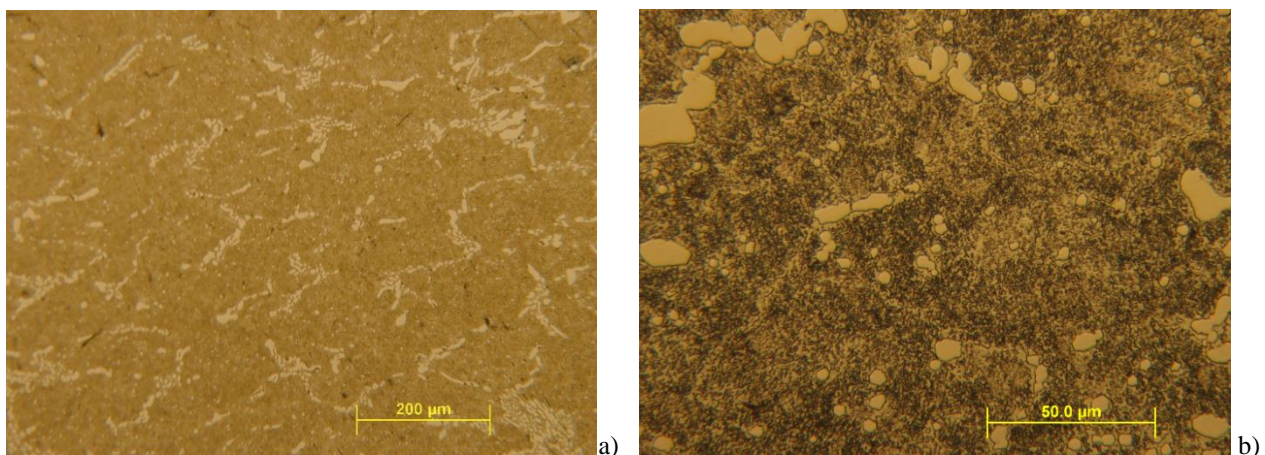


Figure 2. Micrograph of the samples in delivery condition: annealed a)  $N_{original}=100\times$  b)  $N_{original}=500\times$

## Results

In the microstructure of the samples after quenching were observed that the primary carbide did not dissolve at 1070°C during to austenitization and their net structure have not been changed during to heat treatment. The primary austenite grain sizes were ASTM 8. The primary carbides sizes are between 10-50 μm. In etched region small rounded carbides are visible as it showed in *Figures 3*.

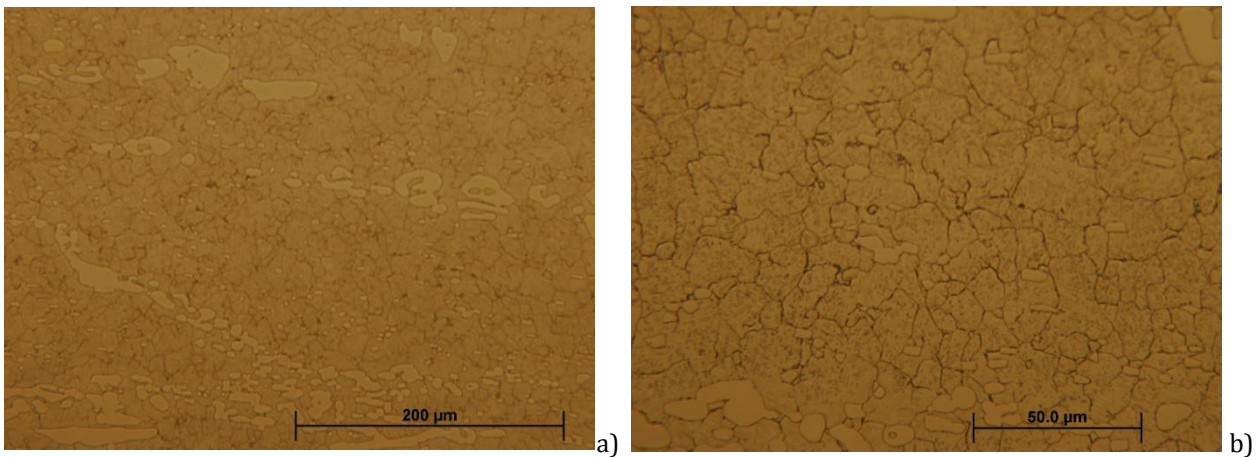


Figure 3. Micrograph of the quenched samples  $N_{original}=100x$  b)  $N_{original}=500x$

The Figure 4 show the microstructure of the samples heat treated conform Figure 1 a). The primary carbide shape and distribution is similar to quenched samples. In the etched structure became visible small compounds precipitations. The hardness of the matrix is 641 HV<sub>1</sub>.

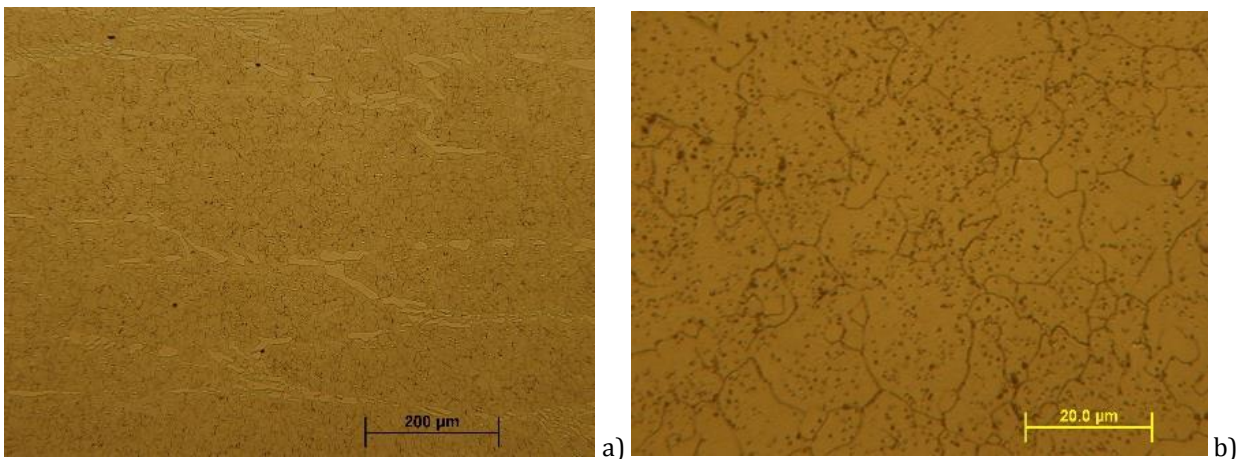


Figure 4. Micrograph of samples quenched and tempered at 200°C. a)  $N_{original}=100x$  b)  $N_{original}=1000x$

After tempering at 510°C the matrix of the samples became homogenous as it presented in Figure 5.a). Studying with higher resolution in matrix can see retained austenite and tempered martensite near small rounded carbides (Figure 5 b).

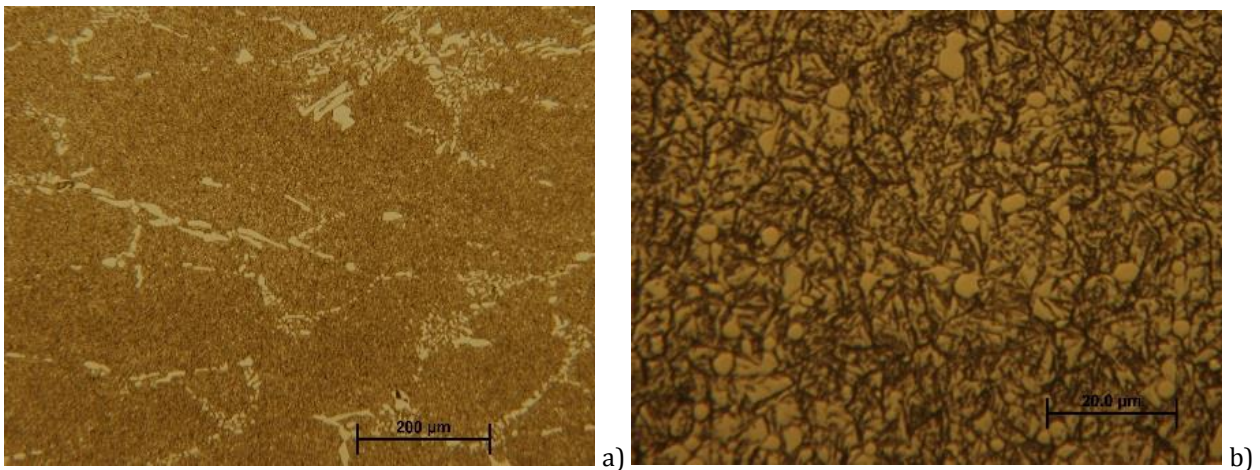


Figure 5. Micrograph of quenched and tempered sample at 510°C. a)  $N_{original}=100x$  b)  $N_{original}=1000x$

The microstructure of a specimen tempered three times is show in Figure 6.

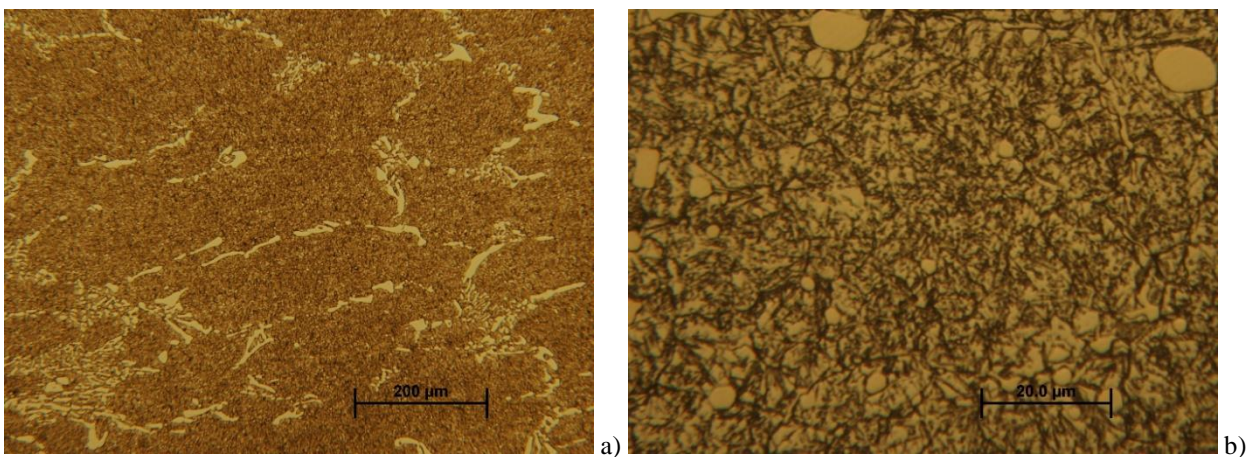


Figure 6. Quenched and triple tempered sample micrographs a)  $N_{original}=100x$  b)  $N_{original}=1000x$

The study the microstructure of the conventionally quenched and triple tempered sample by optical microscope in high resolution show a dense precipitation of carbides and tempered martensite (Figures 6. b). The quantity of retained austenite is reduced compared with the sample tempered one time as it can be seen in Figure 6.b) and Figure 5. b).

The microstructure of cryogenically treated and tempered sample at 510°C is presented in Figure 7. In cryogenically treated sampled retained austenite quantity is decreased comparing with conventional quenched samples like in Figure 5. b) and Figure 7. b). The carbides precipitation are fines and their distribution are homogeneous in etched zones.

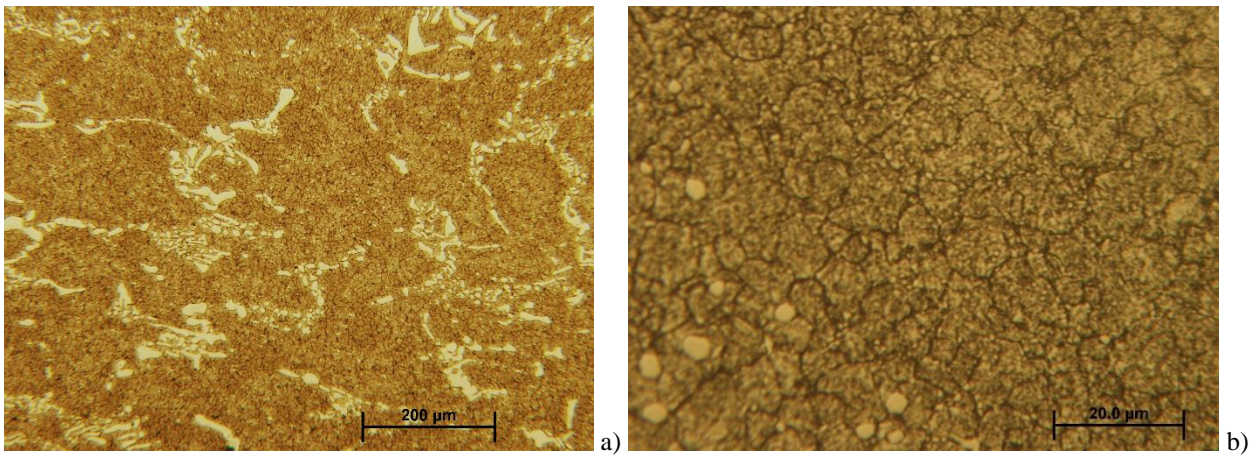


Figure 7. Microstructure of cryogenically treated and tempered sample at 510°C

a)  $N_{original}=100x$  b)  $N_{original}=1000x$

Studying the microstructure of cryogenically treated and three time tempered samples (Figure 8) were observed partially reducing primary carbides sizes and their morphology became more spherical. The distribution of the precipitated carbides is denser and the martensite are full tempered. The retained austenite is minimalized.

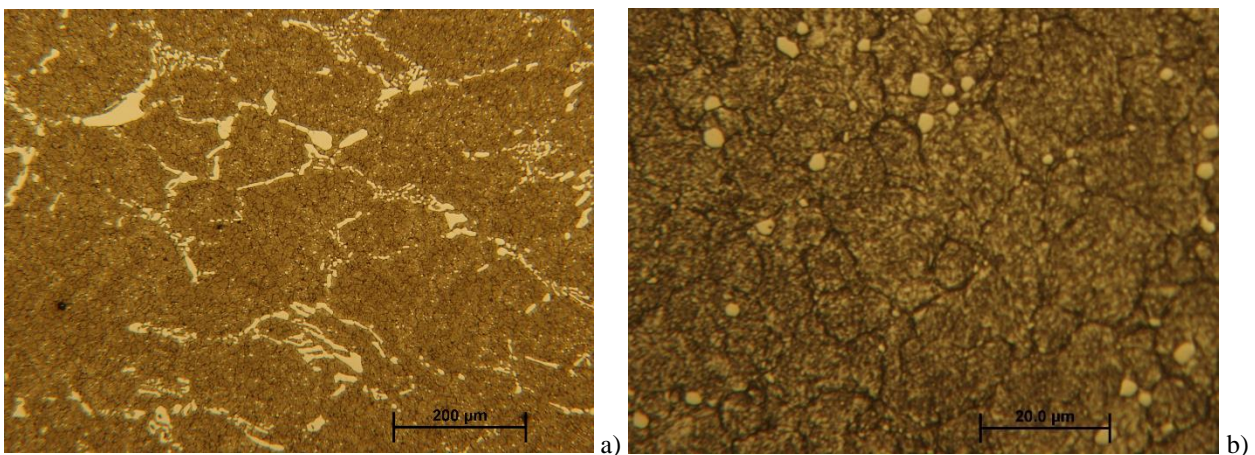


Figure 8. Cryogenically treated and triple tempered sample micrographs. a)  $N_{original}=100x$  b)  $N_{original}=1000x$

The microstructure changings after different heat treatment technologies are reflected in the hardness values. Austenitizing temperatures, cryogenic temperatures tempering temperatures and the hardness evolutions are given in Tab.2.

Studying the microhardness of etched zones were observed that cryogenically treating increase the hardness due to reduction of retained austenite content. In case of conventional quenching increasing the tempering temperature to 510°C the hardness has been increased too. The reason is that the secondary carbides precipitations begins.

After the three times tempering the hardness continues to increase due to the secondary carbides precipitations.

Nr	Austenitizing temperature	Cryogenic treatment	Tempering 1		Tempering 2		Tempering 3	
			T(°C)	HV1	T(°C)	HV1	T(°C)	HV1
1	1070 (°C)	-	200	641	-	-	-	-
2	1070 (°C)	-	510	663	-	-	-	-
3	1070 (°C)	-	510	-	480	-	480	695
4	1070 (°C)	-80	510	746	-	-	-	-
5	1070 (°C)	-80	510	-	480	-	480	738

Table 2. Heat treatment process and hardness evolution

The cryogenic treatment consequence is the higher hardness due to retained austenite reduction.

The best properties were obtained after tempering three times at high temperature.

## Conclusion

In cryogenically treated sampled retained austenite quantity is decreased comparing with conventional quenched samples. Cryogenic treatment accelerate the decomposition of martensite and modify the precipitation behaviour of secondary carbides. The reason for increasing hardness due to the cryogenic treatment is related to the conversion of the retained austenite to martensite and the sediment of the tiny nanometer-sized carbides, and more appropriate distribution of carbides so the transformation of retained austenite to martensite and the secondary carbide precipitation are the main mechanism responsible for the properties improvement of this steel quality.

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