# **Relationship Between Ultrasonic Characteristics and Mechanical Properties of Tempered Martensitic Stainless Steel**

*Cheng-Hsun Hsu, Hwei-Yuan Teng, and Yeong-Jern Chen*

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**This research studied the relationship between the ultrasonic characteristics and the mechanical properties of tempered CA-15 martensitic stainless steel (MSS). The results show that, for as-quenched specimens, a chromium carbide film at the martensitic boundary of the as-cast specimen will disappear causing a change in the mechanical properties (e.g., the tensile strength is decreased or the hardness and the toughness are increased). For the tempered MSS, the correlation of the ultrasonic velocity and the tensile strength, hardness, and toughness is not obvious. However, there is a highly positive correlation with the elastic** modulus  $(E)$  of the material. For the ultrasonic attenuation evaluation, the attenuation coefficient  $(\alpha)$  has **a positive correlation with the tensile strength and the hardness, while there is a negative correlation with the toughness and the elongation. Also, a higher-frequency probe would cause the better sensitivity, but the data are relatively dispersed.**

**Keywords** attenuation coefficient, mechanical property, tempered martensitic stainless steel, ultrasonic velocity

# **1. Introduction**

Martensitic stainless steel (MSS) possesses excellent hightemperature strength and corrosion resistance and is widely used in highly stressed parts, such as turbines and gas distributor heads.<sup>[1]</sup> In general, MSS needs to be heat treated by austenitizing and tempering processes to achieve the desirable mechanical and corrosion properties. Ultrasonic techniques are used not only to detect discontinuities, such as voids, cracks, inclusions, and precipitates, $[2-5]$  but are also used to evaluate such characteristics of a material as microstructure, grain size, and residual stress.<sup>[6-10]</sup> The velocity and attenuation of sound waves are commonly used for such purposes.

The velocity of an ultrasonic wave as it propagates through a solid material is affected by the elastic modulus (*E*), the density ( $\rho$ ), and Poisson's ratio ( $\mu$ ). The relationships among these properties and the longitudinal or transverse velocity of the ultrasonic wave (i.e.,  $V_L$  or  $V_T$ , respectively) can be expressed as follows<sup>[11]</sup>:

$$
V_{\rm L} = \sqrt{\frac{E(1-\mu)}{\rho(1+\mu)(1-2\mu)}}
$$
 (Eq 1)

$$
V_{\rm T} = \sqrt{\frac{E}{2\rho (1 + \mu)}}
$$
 (Eq 2)

**Cheng-Hsun Hsu,** Department of Materials Engineering Tatung University, Taipei, Taiwan 104, Republic of China; and **Hwei-Yuan Teng,** and **Yeong-Jern Chen,** Department of Mechanical Engineering, De Lin Institute of Technology, Tu-Cheng, Taiwan 236, Republic of China. Contact e-mail: chhsu@ttu.edu.tw. **Fig. 1** Heat treatment conditions in this experiment

Equations 1 and 2 imply that the acoustic speed is a constant value and thus is characteristic of a material in a specific state. That is to say, a microstructural change in the material can cause an alteration of the ultrasonic velocity.

When an ultrasonic wave travels through a solid object, its amplitude (*A*) or its intensity (*I*) will decay exponentially with respect to the distance  $(X)$  of the sound propagation.<sup>[11]</sup> The relationship can be written as

$$
A = A_o \exp(-\alpha X), \text{ or } I = I_o \exp(-\alpha X) \qquad \text{(Eq 3)}
$$

where  $A_0$  and  $I_0$  are the initial sound amplitude and the intensity, respectively. The attenuation coefficient  $(\alpha)$  can be further described by $[2,11]$ 

$$
\alpha(f) = \frac{20[\log(A1/A2) + 2 \log R]}{2d}
$$
 (Eq 4)

where  $\alpha(f)$  is in decibels per millimeter, *A*1 and *A*2 are peak amplitudes of the first and second transmitted pulses in millimeters, and *d* is the test specimen thickness in millimeters. The term *, the reflection coefficient of the coupling plane, is equal* to  $(1 - \eta)/(1 + \eta)$ , and  $\eta$  is the acoustic impedance of the



coupling plane. The attenuation coefficient of the ultrasonic wave is affected by absorption (i.e., influenced by dislocation damping, magnetic resistance, and thermal elasticity) and scattering (i.e., influenced by grain boundaries, voids, inclusions, second-phase particles, and cracks).<sup>[8]</sup> The attenuation coefficient is also a function of detector frequency.

It is important to understand the relationships between the ultrasonic characteristics (velocity and attenuation) and the mechanical properties of MSS, tempered at various temperatures

such that the mechanical properties of the material can be evaluated by nondestructive ultrasonic techniques.

# **2. Experimental Procedure**

## *2.1 Material and Heat Treatment*

In this study, 32 mm thick, CA-15-cast MSS plates produced by a regular foundry were used as the experimental material. The chemical composition (wt.%) of the material was



**Fig. 2** SEM and EPMA scan of the experimental material: **(a)** as-cast; **(b)** as-quenched; **(c)** tempered at 400 °C; and **(d)** tempered at 600 °C



**Fig. 3** Effect of the tempering treatment on the mechanical properties of the experimental material: **(a)** tensile strength and elongation; and **(b)** hardness and impact toughness





analyzed by emission spectroscopy as follows: Fe-0.14 C-0.34 Si-0.33 Mn-12.1 Cr-1.02 Ni-0.017 P-0.001 S. This chemistry conformed to the CA-15 nominal chemical specifications. Various specimens for mechanical testing and ultrasonic evaluation were machined from the cast plates and then were heat treated to different schedules. The heat treatment procedure involved austenitizing, quenching, and tempering, as shown in Fig. 1.

#### *2.2 Mechanical Testing and Microstructural Analysis*

Hardness testing of the samples was performed using a Vickers hardness tester (30 kg load). Five hardness readings were taken and averaged for each specimen. Tensile tests were carried out using a dynamic testing machine (MTS model 810 MTS Systems Corporation, MN). Specimens for tensile testing were machined according to the subsize tensile specimen of the ASTM E 8M standard.<sup>[12]</sup> Impact tests were carried out using a Charpy impact tester. Specimens for impact testing were machined according to the V-notch impact specimen of the ASTM E 23 standard.<sup>[13]</sup> Three tests were performed, and the results were averaged to represent the tensile strength, elongation, and impact toughness of the experimental material in each heat-treated condition.

Scanning electron microscopy (SEM), electron probe microanalysis (EPMA), and transmission electron microscopy (TEM) were used to examine the microstructure of the material. The specimens were polished and etched with Vilella's reagent (5 mL HCl + 1 g picric + 100 mL ethanol) for SEM and EPMA observations.

#### *2.3 Ultrasonic Measurement*

An ultrasonic A-scan pulsing instrument was attached to a gate monitor (model USIP12 with DTM 12 attachment, Krautkramer, Branson, PA) for ultrasonic measurement of the velocity, and for the first and second pulse peak amplitudes (i.e., the values *A*1 and *A*2 used in Eq 4) for attenuation coefficient calculations. Ultrasonic probes with frequencies of 1 and 5 MHz were used, and commercial motor oil was adopted as the coupling agent for the ultrasonic contact-type testing in these experiments. A longitudinal wave was propagated through the sample to evaluate the acoustic characteristics at the same position before and after heat treatments. Five ultrasonic readings of each specimen were taken and averaged to represent the data obtained.

# **3. Results and Discussion**

## *3.1 Microstructure and Mechanical Properties*

Microstructure of the experimental as-cast material mainly consisted of martensite and ferrite phases in the matrix along with chromium carbide films around grain boundaries (Fig. 2a). The carbide films were effectively eliminated by austenitizing at 1010 °C for 4 h followed by quenching, as shown in Fig. 2(b). Figure 2(c) shows the micrograph of the sample tempered at 400 °C. The microstructure was obviously unaltered, but the Cr content at the grain boundary of the martensitic phase had increased. Furthermore, ferrite islands started to precipitate in the martensite grains. After tempering at 500 and 600 °C, Cr and C were found to be uniformly dispersed in the matrix. Figure 2(d) shows micrographs of the sample tempered at 600 °C, at which point ferrite islands have precipitated in the martensite grains.

The mechanical properties of the material after different tempering treatments are listed in Table 1 and are illustrated in Fig. 3. After austenitization and quenching, the hardness and impact toughness were higher than those of the as-cast sample, but tensile strength was lower. The reason is that Cr carbides dissolve in the matrix for the austenitized specimens causing a hardness increase in the material. The increase of impact toughness resulted from the disappearance of boundary carbides, which could dominate the intergranular fracture in the matrix. However, due to the boundary carbide effect and the retarded microcrack propagation being nonexistent, a reduction in tensile strength resulted. After tempering at 300-400 °C, secondary hardening/strengthening occurred for the material as a result of tiny carbides precipitating at the grain boundaries. These carbides can be seen clearly in the TEM micrographs (Fig. 4). This caused tempering mar-



Carbides in 403 b<br>3U0112 200.OKV X50K 100nm **Fig. 4** TEM micrographs of the sample tempered at 400 °C: **(a)**





tensite embrittlement (TME) and led to a reduction in impact toughness. After 500 and 600 °C tempering, the hardness and tensile strength of the material decreased, while the impact toughness increased, as a result of the softening of the matrix.

## *3.2 Interrelation of Ultrasonic Velocity and Mechanical Properties*

The ultrasonic characteristics from various tempered specimens are listed in Table 2. Figure 5(a-d) illustrates the relationship between ultrasonic velocity and mechanical properties. From these figures, it is clearly seen that the data were quite scattered and that no particular relationship exists among ultrasonic velocity hardness, toughness, or tensile strength. However, it is interesting that there was a distinct positive correlation (correlation coefficient >0.85) between the ultrasound velocity and the *E* of the specimens (except for the as-cast sample). These results are shown in Fig. 6. The *E* of the material was changed by tempering treatments, and the ultrasonic velocity increased with an increase in *E*. Also, the as-cast sample possesses a higher value of *E* than that of the tempered samples and results from the carbide films at the grain boundaries retarding the elastic deformation of the material in the linear elastic strain region.

Incidentally, for detection with the 1 MHz probe, the variation of ultrasonic velocity was similar to that of the 5 MHz probe. However, the ultrasonic velocity at 1 MHz was higher than that at 5 MHz. The reason for this is yet unknown.

## *3.3 Interrelation of Ultrasonic Attenuation and Mechanical Properties*

Figure 7 depicts the relationship between the ultrasonic attenuation coefficient  $\alpha$  and the mechanical properties. The ascast specimen showed substantial acoustic attenuation compared with the heat-treated specimens, and this might have resulted from the carbide films at the grain boundaries causing a scattering of the acoustic wave. Therefore, despite the anomalous behavior of the as-cast specimen, linear equations were derived and are shown in Fig. 7. It can be clearly seen that strength and hardness are positively correlated with the attenu-

beaded boundary carbide; and **(b)** stringed carbide



**Fig. 5** Relationship between the ultrasonic velocity and the mechanical properties of the experimental materials, under different frequencies: **(a)** hardness; **(b)** tensile strength; **(c)** elongation; and **(d)** impact toughness

ation coefficient, while toughness and elongation are negatively correlated with it. Correlation coefficients of all the linear equations were >0.8. This correlation implies that the attenuation coefficient plays an important role in evaluating the mechanical properties of the material. Table 3 shows the correlation equations for relationships between the ultrasonic attenuation coefficient and various mechanical properties and provided the formula for the evaluation of mechanical properties based on the nondestructive ultrasonic attenuation data. In addition, the higher the ultrasonic frequency, the higher the attenuation coefficient obtained. However, it also resulted in greater scatter of the attenuation data.

# **4. Conclusions**

The purpose of this research was to derive relationships among the ultrasonic characteristics and mechanical properties of tempered CA-15 MSS. The following conclusions can be made:



**Fig. 6** Relationship between the ultrasonic velocity and the *E* of the experimental materials, under different frequencies (*r* is the correlation coefficient)



**Fig. 7** Relationship between the ultrasonic attenuation and the mechanical properties of the materials, under different frequencies: **(a)** hardness; **(b)** tensile strength; **(c)** elongation; and **(d)** impact toughness (*r* is the correlation coefficient)

**Table 3 Correlation equations of attenuation coefficient and mechanical properties using different frequencies**

<b>Probe frequency</b>	1 MHz	5 MHz
Hardness, HV	$Y = 17.401X - 552$	$Y = 3757X - 157$
Tensile strength, MPa	$Y = 64,277X - 2289$	$Y = 12,039X - 600$
Elongation, %	$Y = -4156X + 2198$	$Y = -680X + 97$
Impact toughness, J	$Y = -3788X + 204$	$Y = -893X + 128$

Note: *X,* attenuation coefficient; *Y,* mechanical properties

- The microstructure of the material consisted mainly of martensite and ferrite with carbide films around the martensite interphase boundaries. These carbide films retarded microvoid propagation (i.e., increased the strength) during tensile testing but caused grain boundary embrittlement (i.e., reduced the toughness) during impact testing and distinctly scattered and attenuated the propagated acoustic waves.
- The ultrasonic velocity of the wave in the samples, except for the as-cast material, was positively correlated with the *E*. The relationship between the ultrasonic velocity and the mechanical properties was not obvious.
- The attenuation coefficient was positively correlated with the strength and the hardness of the material, but was negatively correlated with the toughness and elongation.
- For ultrasonic characteristic evaluation, the higher frequency (5 MHz) probe, compared with the 1 MHz probe, had better sensitivity, a higher attenuation coefficient  $(\alpha)$ , and a lower velocity value. However, using this probe resulted in relatively dispersed data.

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