

Fig. 3. Effect of oxygen pressure on the oxidation of Ferrovac E iron at 350°C.

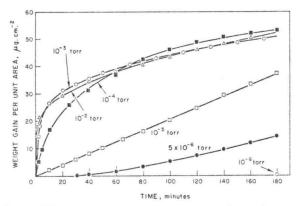


Fig. 4. Effect of oxygen pressure on the oxidation of Ferrovac E iron at 400 $^\circ\text{C}.$

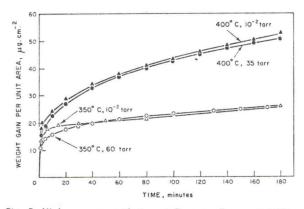


Fig. 5. High pressure oxidation of Ferrovac E iron at 350° and 400°C.

creases with increasing oxygen pressure in the range 10^{-4} to 35 or 60 Torr of oxygen, the weight gain after 3 hr is essentially pressure independent, as summarized in Table I. After 3 hr oxidation at 350°C, the average weight gain for this pressure range is about 26 μ g

Table I. Weight gain per unit area after 3 hr oxidation

	Weight gain (µg cm-2) after 3 hr	
	(a)	(b)
Pressure, Torr	350°C	400°C
10-8	~0.5	~0.1
5×10^{-6}	17.0	14.2
10-5	22.6	37.0
10-4	24.8	53.2
10-3	27.3	51.0
10-2	25.8	53.0
35		50.9
60	25.1, 25.0, 25.8	

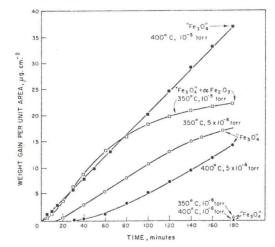


Fig. 6. Comparison of the low pressure oxidation of Ferrovac E iron at 350° and 400°C.

cm⁻², compared with a value of about 52 μg cm⁻² at 400°C. An induction period is observed for oxidations at 5 x 10⁻⁶ Torr at both 350° and 400°C.

The weight gains after 3 hr of oxidation at 10^{-6} Torr were obtained by difference from the weight of the specimen before and after oxidation, as determined by the Mettler analytical microbalance, (taking into account the removal of the oxide film resulting from the electropolishing), the Cahn electrobalance being not sufficiently stable to identify this small weight change.

The kinetic data from oxidations at 10^{-4} , 10^{-3} , and 10^{-2} Torr oxygen at 350° and 400°C is plotted in a parabolic manner in Fig. 7 and 8, respectively. Another parabolic plot of the data from the oxidation at 35

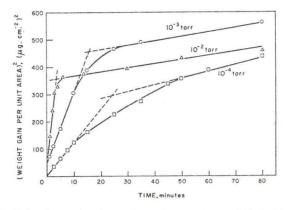


Fig. 7. Parabolic plot of data from the oxidation at 350°C; 10^{-4} , 10^{-3} , and 10^{-2} Torr oxygen pressure.

