

INTERACTION BETWEEN ACTIVE PLASMA AND GROWING Co-C-O – LAYER DURING PACVD

Arnold Nürnberg, Ralf Stolle, Georg Wahl and Karlo T. Raic*

Institut für Oberflächentechnik und Plasmatechnische Werkstoffentwicklung, TU Braunschweig, Bienroder Weg, 53, D-38108 Braunschweig, Germany,

*Faculty of Technology and Metallurgy, Belgrade University, Karnegijeva 4, POB 3503, 11120 Belgrade, Yugoslavia / Serbia

Transport and chemical rate phenomena are analysed for Co-C-O-layers generated on steel C35 from cobalt(II) acetylacetonate by PACVD. The investigations are carried out in a parallel plate reactor, under the high frequency (13.56 MHz) plasma, with low ionization (ionization degree 10^{-7} - 10^{-6} , electron temperature 5 eV, electron density $0.6 \cdot 10^{15}$ to $1.8 \cdot 10^{15} \text{ m}^{-3}$) and at the pressure of 100 – 400 Pa, see Figure 1.

Deposition rate phenomena are estimated according to the semi-theoretical concept based on laminar boundary layer theory. On the other side, in active plasma, the concept of steady state diffusion with concentration source/sink is used to substitute the net deviation of characteristic concentration flow in plasma above the substrate.

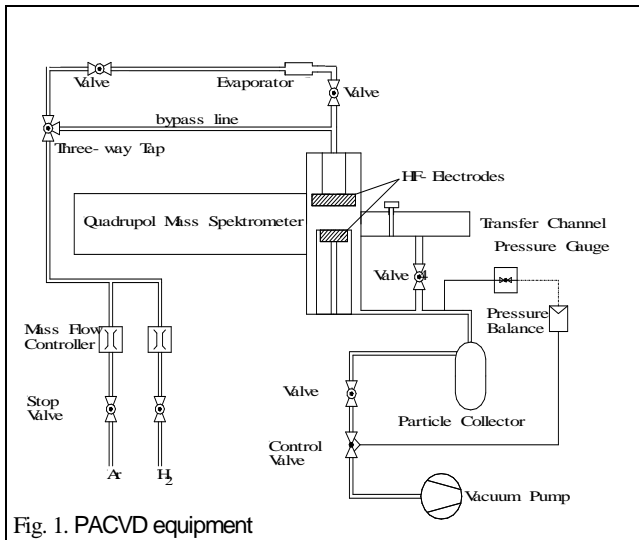


Fig. 1. PACVD equipment

Table 1 Calculated values

	523 K	623 K	723 K
Re	1.89	1.67	1.50
Sc	1.12	1.16	1.18
Sh	1.42	1.36	1.29
δ_v , (mm)	35.6	37.9	40.0
δ_c , (mm)	34.3	36.0	38.0
D_{100Pa} (m^2/s)	$6.73 \cdot 10^{-2}$	$8.86 \cdot 10^{-2}$	$11.1 \cdot 10^{-2}$
D_{200Pa} (m^2/s)	$3.36 \cdot 10^{-2}$	$4.43 \cdot 10^{-2}$	$5.55 \cdot 10^{-2}$
D_{400Pa} (m^2/s)	$1.68 \cdot 10^{-2}$	$2.21 \cdot 10^{-2}$	$2.77 \cdot 10^{-2}$
$D_{10^{-5}Pa}$ (m^2/s)	$6.73 \cdot 10^{-5}$	$8.86 \cdot 10^{-5}$	$11.1 \cdot 10^{-5}$
β_{100Pa} (m/s)	1.27	1.60	1.90
β_{200Pa} (m/s)	0.63	0.80	0.95
β_{400Pa} (m/s)	0.31	0.40	0.47
$\beta_{10^{-5}Pa}$ (m^2/s)	$1.27 \cdot 10^{-3}$	$1.60 \cdot 10^{-3}$	$1.90 \cdot 10^{-3}$

In the system Co-C-O the deposition is strongly influenced by a particle formation in the gas phase.

In spite of the vacuum sealed equipment the layers always contain oxygen. Therefore the oxygen comes from the precursor compounds. The deposition temperature is rather low, the maximum temperature is 750 K. Because of these reasons, the PACVD deposition of Co-C-O layers is discussed in the light of transport and chemical rate phenomena at low pressures with plasma activated chemical reactions in gas phase.

MODELING About the semi-theoretical approach based on laminar boundary layer theory see (8-10). According to balance equation, referent situation, simplified analysis of Frank-Kamenetskii (13) and the model of mass transfer coefficient (β), Table 1, one can obtain deposition rate r (Table 2.)

$$j = \beta \theta^{\text{III}} \Delta C / [1 + (z\Phi)^{-1}] = (\rho/M) d\delta_r/dt \quad \text{or}$$

$$r = d\delta_r/dt = (M/\rho) \beta \theta^{\text{III}} \Delta C / [1 + (z\Phi)^{-1}]$$

where: $\Phi = [1 - (\pm f(m)_V \pm f(m)_D \pm FG_H)/N_V]$, numerical value of real conditions, $z = k(s)/\beta$, ρ : the density of Co, M : the mass of Co per molecule and θ^{III} : normalized deviation, derived from the concept of steady state diffusion with concentration source/sink in surrounding plasma (14).

Table 2. Values for ΔC and (r)

mol/m^3 ($\text{mol/m}^2\text{s}$)	523 K	623 K	723 K
$\Delta C_{100Pa} \cdot 10^{-6}$ ($r \cdot 10^{-6}$)	3.1 (≈ 4.0)	4.3 (≈ 7.0)	4.7 (≈ 9.0)
$\Delta C_{200Pa} \cdot 10^{-6}$ ($r \cdot 10^{-6}$)	12.7 (≈ 8.0)	11.2 (≈ 9.0)	12.7 (≈ 12.0)
$\Delta C_{400Pa} \cdot 10^{-6}$ ($r \cdot 10^{-6}$)	11.1 (≈ 3.5)	13.7 (≈ 5.5)	21.0 (≈ 10.0)

The presented procedure is imagined as a flexible iterative system for the following experimental data. This model can be used as a tool in computer-aided process optimization, too.

REFERENCES

1. Wiesemann, *Pure & Appl. Chem.*, **68**, 1029(1996)
2. Rhee, S., Szekeley, J., *J. Electrochem. Soc.: Solid-State Science and Technology*, **133**, 2194(1986)
3. Kushner, M. J., *J. Appl. Phys.*, **8**, 63 (1988)
4. Gladush, G. G., Rodionov, N. B., *J. Chem. Vap. Deposition*, **6**, 76(1997)
5. Jablonovski, J., *Cobalt*, **14**, 28 (1962)
6. Charles, R.G. and Haverlack, P.G., *J. Inorg. Nucl. Chem.*, **31**, 995 (1969)
7. Gu, S. et al, *Thin Solid Films*, **45**, 340 (1999)
8. Raic, K.T., *ISIJ Int.*, **32**, 514 (1992).
9. Raic, K.T., *J. de Physique IV*, C5, **5**, 235 (1995).
10. Raic, K.T., *Interface Science*, **8**, 85 (2000)
11. Schlichting, H., *Boundary-Layer Theory*, Pergamon Press, New York, (1955).
12. Bird, R.B, Stewart W.E. and Lightfoot, E.D., *Transport Phenomena*, John Wiley and Sons, New York, (1960).
13. Frank-Kamenetskii, D.A., *Diffusion and Heat Exchange in Chemical Kinetics*, Princeton University Press, Princeton, (1960).
14. K.T.Raic, *Surf.&Coat.Techn.*, **92**, 22 (1997)
15. Reid, R.C., Prausnitz, J.M. and Poling, B.E., *The Properties of gases and liquids*, McGraw-Hill Book Co., New York, 4th ed., (1987)

