

# Reactive High Power Impulse Magnetron Sputtering (HiPIMS)

Jón Tómas Guðmundsson<sup>1</sup>, Friðrik Magnus<sup>1,2</sup>,  
Tryggvi K. Tryggvason<sup>1,3</sup>, Ólafur B. Sveinsson<sup>1,4</sup>,  
S. Shayestehaminzadeh<sup>1</sup>, and Sveinn Ólafsson<sup>1</sup>

<sup>1</sup> Science Institute, University of Iceland, Iceland

<sup>2</sup> Uppsala University, Sweden

<sup>3</sup> Lund University, Sweden

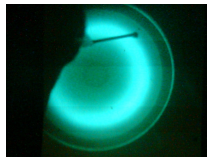
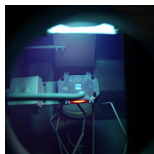
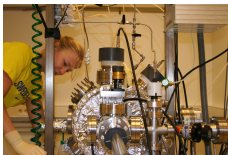
<sup>4</sup> Linköping University, Sweden

tumi@hi.is

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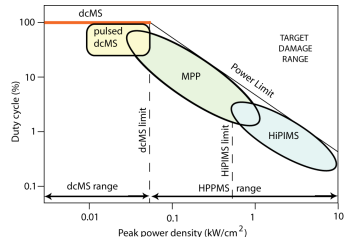
# Introduction



- Magnetron sputtering has been the workhorse of plasma based sputtering methods for over three decades
- For many applications a high degree of ionization of the sputtered vapor is desired
- In a HiPIMS discharge a high power pulse is supplied for a short period
  - low frequency
  - low duty cycle
  - low average power
- Ionized flux of sputtered vapor introduces an additional control parameter into the deposition process

# Introduction

- High power pulsed magnetron sputtering (HPPMS)
- HiPIMS
  - a pulse of very high amplitude, an impulse, is applied to the cathode and a long pause exists between the pulses
- Modulated pulse power (MPP)
  - the initial stages of the pulse (few hundred  $\mu\text{s}$ ) the power level is moderate (typical for a dcMS) followed by a high power pulse (few hundred  $\mu\text{s}$  up to a ms)



From Gudmundsson et al. (2012), JVSTA **30** 030801

- Power density limits
  - $p_t = 0.05 \text{ kW}/\text{cm}^2$  dcMS limit
  - $p_t = 0.5 \text{ kW}/\text{cm}^2$  HiPIMS limit

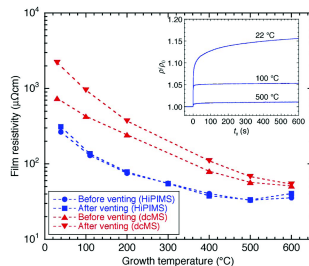
# Reactive HiPIMS - Applications





## Application – Film Resistivity

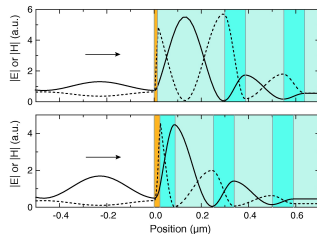
- TiN as diffusion barriers for interconnects
- HiPIMS deposited films have significantly lower resistivity than dcMS deposited films on  $\text{SiO}_2$  at all growth temperatures due to reduced grain boundary scattering
- Thus, ultrathin continuous TiN films with superior electrical characteristics and high resistance towards oxidation can be obtained with HiPIMS at reduced temperatures



From Magnus et al. (2012) IEEE EDL **33** 1045

## Application – Bragg mirror

- Multilayer structures containing a high-contrast ( $\text{TiO}_2/\text{SiO}_2$ ) Bragg mirror fabricated on fused-silica substrates
  - reactive HiPIMS  $\text{TiO}_2$  (88 nm)
  - reactive dcMS  $\text{SiO}_2$  (163 nm)
  - capped with semitransparent gold
- Rutile  $\text{TiO}_2$  ( $n = 2.59$ ) and  $\text{SiO}_2$  ( $n = 1.45$ ) provide a large index contrast
- Smooth rutile  $\text{TiO}_2$  films can be obtained by HiPIMS at relatively low growth temperatures, without post-annealing



From Leosson et al. (2012) Opt. Lett. **37** 4026

Magnus et al. (2011) Mater. Res. Soc. Symp. Proc. Vol. 1352

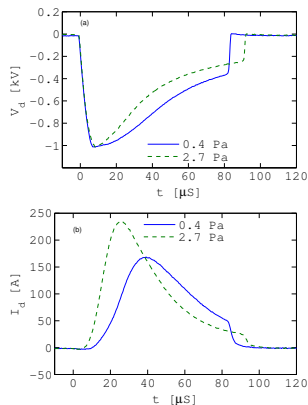
Agnarsson et al. (2013) TSF **545** 445



# Reactive HiPIMS - Voltage - Current - Time characteristics

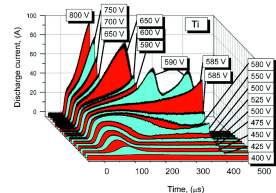
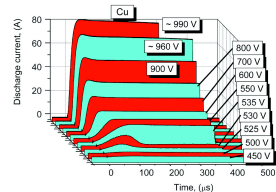
# HiPIMS - Voltage - Current - time

- To describe the discharge current-voltage characteristics the current-voltage-time space is required
- The early work on HiPIMS used 50 – 100  $\mu\text{s}$  pulses and a pulse repetition frequency in the range 50–1000 Hz
- The cathode voltage and the discharge current depend on the discharge gas pressure



# HiPIMS - Voltage - Current - time

- For longer pulses the initial pressure dependent current peak is followed by a second phase that is power and material dependent
- The initial phase is dominated by gas ions, whereas the later phase has a strong contribution from self-sputtering
- For some materials, the discharge switches into a mode of **sustained self-sputtering**

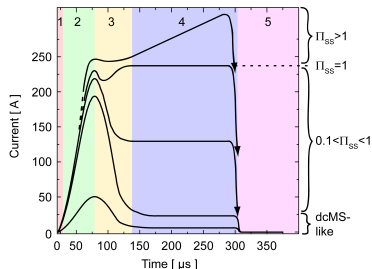


From Anders et al. (2007),

JAP 102 113303 and JAP 103 039901

# HiPIMS - Voltage - Current - time

- A schematic illustration of the discharge current assuming square shaped voltage pulses
- The current is generally characterized by an initial peak followed by a more or less stable current plateau (bottom current curves)
- In other cases it shows an initial peak followed by a second increase of the discharge current (top current curves)



From Gudmundsson et al. (2012), JVSTA **30** 030801

## *HiPIMS - Voltage - Current - time*

- The self-sputtering can operate in a self-sustained mode, when the ions of the sputtered vapor are created at high enough rate that the ions of the working gas are not needed
- The condition for sustained self-sputtering is expressed as

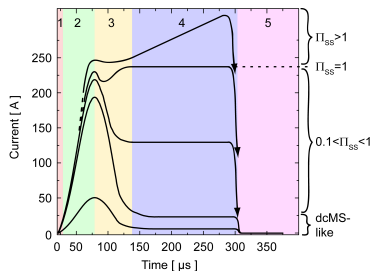
$$\Pi_{ss} = \alpha \beta_t Y_{ss} = 1$$

where

- $\alpha$  is the probability of ionization of the sputtered atom
- $\beta_t$  is the probability that the newly formed ion of the sputtered vapor returns to the target
- $Y_{ss}$  is the self-sputter yield of the ion
- This is a steady state situation and the current remains constant

# HiPIMS - Voltage - Current - time

- The bottom curve represents a range of low self-sputtering,  $\Pi_{ss} < 0.1$  and the discharge physics in the plateau/runaway phase is dcMS-like
- The middle range of power densities, with  $0.1 < \Pi_{ss} < 1$ , represents partially self-sputtering discharge
- The top curve represents self-sputtering runaway which requires  $\Pi_{ss} > 1$  and a self-sputter yield  $Y_{ss} > 1/(\alpha\beta_t) > 1$

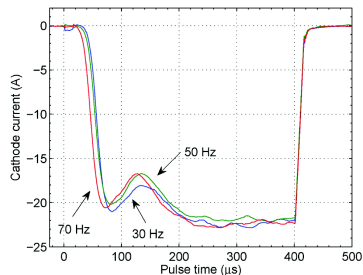


From Gudmundsson et al. (2012), JVSTA **30** 030801



# HiPIMS - Voltage - Current - time

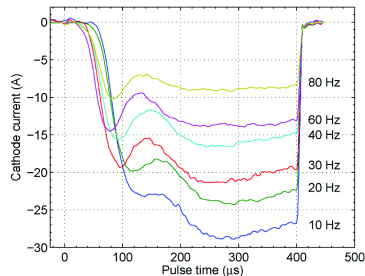
- Ar discharge with Ti target
- The initial peak in current results large flux of atoms from the target
- Collisions of the sputtered atoms with the working gas result in heating and expansion of the working gas – **rarefaction**
- A significant fraction of the sputtered atoms experience electron impact ionization (the ionization mean free path  $\sim 1$  cm) and are attracted back to the target to participate in the sputtering process – **self-sputtering**



From Magnus et al. (2011) JAP **110** 083306

# HiPIMS - Voltage - Current - time

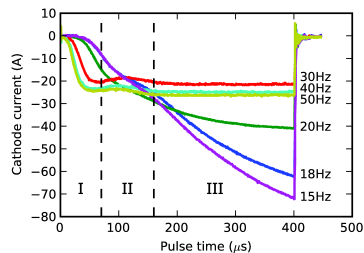
- During reactive sputtering, a reactive gas is added to the inert working gas
- The current waveform in the reactive Ar/N<sub>2</sub> HiPIMS discharge is highly dependent on the pulse repetition frequency, unlike for pure Ar
- N<sub>2</sub> addition changes the plasma composition and the target condition can also change due to the formation of a compound on its surface



From Magnus et al. (2011) JAP **110** 083306

## HiPIMS - Voltage - Current - time

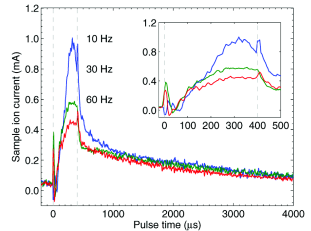
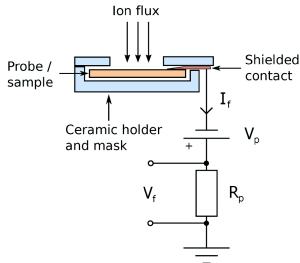
- Similarly for the Ar/O<sub>2</sub> discharge, the current waveform is highly dependent on the repetition frequency and applied voltage which is linked to oxide formation on the target
- The current is found to increase significantly as the frequency is lowered



From Magnus et al. (2012), JVSTA **30** 050601

# HiPIMS - Voltage - Current - time

- The observed changes in the discharge current are reflected in the flux of ions impinging on the substrate



From Magnus et al. (2011), JAP **110** 083306

## HiPIMS - Voltage - Current - time

- The discharge current  $I_d$  is the sum of the ion current  $I_i$  and the secondary electron current  $I_i\gamma_{SE}$  or

$$I_d = I_i(1 + \gamma_{SE})$$

where  $\gamma_{SE}$  is the secondary electron emission coefficient of the target material

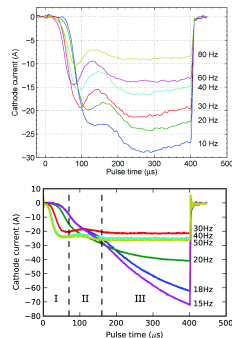
- Also

$$I_i \propto n_i \propto \frac{1}{\mathcal{E}_T}$$

- The total energy loss per electron-ion pair lost from the system  $\mathcal{E}_T$  is expected to increase with the addition of nitrogen
- We must turn to the secondary electron emission yield to explain the self-sputtering runaway and observed frequency dependence of the current in the reactive discharge

# HiPIMS - Voltage - Current - time

- HiPIMS differs significantly from dcMS, due to the fact that self-sputtering quickly becomes dominant and the working gas ions (mostly  $\text{Ar}^+$  and  $\text{N}_2^+$  or  $\text{O}_2^+$ ) are depleted from the area in front of the target, due to rarefaction
- The secondary electron emission yield is governed by the composition of the target (Ti or TiN or  $\text{TiO}_2$ ) and the type of ions that are bombarding it

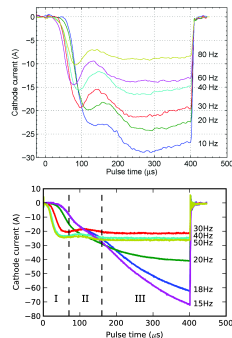


From Magnus et al. (2011), JAP **110** 083306

and Magnus et al. (2012), JVSTA **30** 050601

# HiPIMS - Voltage - Current - time

- $\gamma_{SE}$  is practically zero for singly charged metal ions impacting a target of the same metal
- $\gamma_{SE}$  will be higher for self sputtering from a TiN or TiO<sub>2</sub> target, where N<sup>+</sup>-ions or O<sup>+</sup>-ions are also present, than for self-sputtering from a Ti target, where multiply charged Ti ions are needed to create secondary electrons

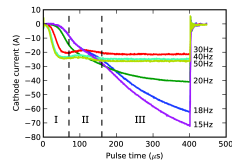
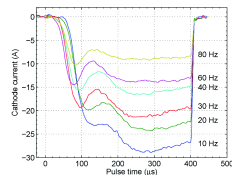


From Magnus et al. (2011), JAP **110** 083306

and Magnus et al. (2012), JVSTA **30** 050601

# HiPIMS - Voltage - Current - time

- At high frequencies, nitride or oxide is not able to form between pulses, and self-sputtering by  $\text{Ti}^+$ -ions (singly and multiply charged) from a Ti target is the dominant process
- At low frequency, the long off-time results in a nitride or oxide layer being formed on the target surface and self-sputtering by  $\text{Ti}^+$ - and  $\text{N}^+$ -ions or  $\text{O}^+$ -ions from  $\text{TiN}$  or  $\text{TiO}_2$  takes place



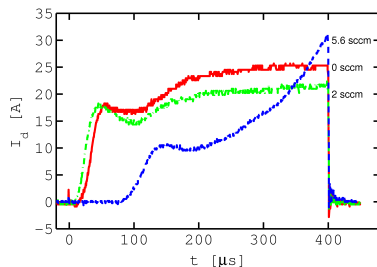
From Magnus et al. (2011), JAP **110** 083306

and Magnus et al. (2012), JVSTA **30** 050601



# HiPIMS - Voltage - Current - time

- As the oxygen flow is increased a transition to oxide mode is observed – The delay in the onset of the current increases, the initial current peak is lowered and a transition to a self-sputtering runaway occurs
- It has been confirmed that in the oxide mode, the discharge is dominated by  $O^+$ -ions, due to oxygen atoms sputtered off the target surface



The current waveforms for an Ar/O<sub>2</sub> discharge with a Ti target where the oxygen flow rate is varied – 600 V, 50 Hz and 0.6 Pa

# Summary



## Summary

- The current-voltage-time waveforms in a reactive discharge exhibit similar general characteristics as the non-reactive case
  - the current rises to a peak, then decays because of rarefaction before rising to a self-sputtering dominated phase
- At low repetition frequency, the long off-time results in a nitride or oxide layer being formed on the target surface and self-sputtering by  $\text{Ti}^+$  and  $\text{N}^+$  or  $\text{O}^+$ -ions from  $\text{TiN}$  or  $\text{TiO}_2$  takes place with an increase in secondary electron emission yield and a corresponding increase in discharge current



# References

The slides can be downloaded at

<http://langmuir.raunvis.hi.is/~tumi/hipims.html>

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