PREDICTIVE THERMAL MODEL FOR INDIRECT TEMPERATURE MEASUREMENT INSIDE ATOMIC CELL OF NUCLEAR MAGNETIC RESONANCE GYROSCOPE

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ABSTRACT

This paper describes a methodology for predicting temperatures inside a micro-device at critical locations where thermal sensors cannot be placed. The device under study is a nuclear magnetic resonance gyroscope, an instrument that is highly sensitive to temperature variations. A physical thermal model of the device is developed and validated using experimental measurements from a prototype. A compact thermal model which approximates the behavior of the physical system is obtained through a sequential selection algorithm (with RMS error of approximately 0.15°C). The compact thermal model can be solved rapidly and is therefore suitable for real-time feedback control of the temperature inside the atomic cell.

KEYWORDS

Predictive models, atomic gyroscope, nuclear magnetic resonance, atomic MEMS, compact thermal model.

INTRODUCTION

We present a two-step process for predicting temperatures inside a device at critical locations where high accuracy is required and direct temperature measurements are not feasible due to generation of parasitic magnetic fields by a temperature sensor, limited space inside the device, or inaccessibility of the area of interest. The methodology described in this paper can be applied to a broad class of atomic MEMS devices. Conventional approaches are effectively open-loop, and do not yield sufficient precision demanded by recent emerging atomic MEMS applications [1].

A Finite Element (FE) model of the device is directly correlated to experimental measurements by matching and using an optimization algorithm. We then develop a Compact Thermal Model (CTM) which reproduces the results of the FE model, with high accuracy, at the critical locations of interest inside the device, such as inside of the atomic cell.

The NMR gyroscope [2-3] is a micromachined sensor which measures inertial rotation by detecting the NMR frequency shift in hyperpolarized noble gases (Fig. 1). A circularly polarized laser beam travels through optically pumped alkali vapor and noble gases confined in a cell. The laser frequency is tuned to the resonant absorption frequency of the alkali isotope. The light transmitted through the cell is collected on a photodetector, and inertial rotation is measured by detecting a shift in the

nuclear precession frequency, which is embedded in the optical signal due to spin-exchange between the nuclei and the alkali vapor [4]. The degree of polarization is highly dependent on the cell temperature, and temperature fluctuations in the cell can therefore create undesirable shifts in the signal. The temperature sensitivity of an NMR gyroscope is theoretically estimated to be on the order of 1 deg/h/mK [5-6]. Therefore the temperature stability of the cell is required to be better than 1 mK to achieve the targeted sub-degree per hour gyroscope performance.

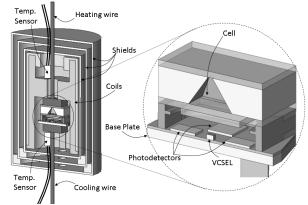


Figure 1: Location of components and temperature sensors in the NMR gyroscope.

High vapor pressure of the confined alkali is needed to achieve a sufficiently strong absorption signal. ⁸⁷Rb isotope is chosen for its high vapor pressure at relatively low temperatures. A cell temperature of 120°C is chosen to achieve high polarization with uniform distribution in the cell [6]. In addition, the light source, a Vertical-Cavity Surface-Emitting Laser (VCSEL), must operate at temperatures below 80°C to avoid degradation and reduced lifetime. Due to the close proximity of the cell and the VCSEL, active heating and cooling was included in the presented prototype through an external heater and a ThermoElectric Cooler (TEC) (Fig. 1). The cell and VCSEL temperatures need to be measured and controlled to ensure that they satisfy the stringent thermal requirements.

EXPERIMENTAL MEASUREMENTS

An NMR gyroscope prototype was built, Fig. 1, and thermal experiments conducted on the setup shown in Fig. 2. The NMR gyroscope prototype and signal conditioning electronics were enclosed in a metallic box to reduce noise in the measurements. A DAQ board from National Instruments was used to acquire the gyroscope signal and

temperature measurements. Heater and TEC controllers (TC200 and TED200C from Thorlabs) were used to tune the heating and cooling temperatures. Function generators were used to sweep the VCSEL current and to supply the signals for magnetic field generation in the gyroscope.

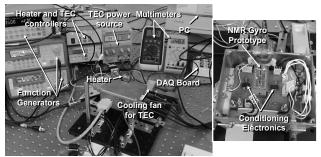


Figure 2: Experimental setup for thermal characterization of the NMR Gyroscope.

Temperature Measurements

Two temperature sensors (Thorlabs TH100PT) were placed inside of the device, but outside of the cell (Fig. 1). The first sensor provided temperature measurements a few millimeters above the cell. The second sensor provided measurements below the base plate and near the VCSEL. Two additional temperature sensors (Thorlabs TH10K) were used to measure the heater and TEC temperatures. In addition to these four direct temperature measurements, two indirect temperature measurements, the VCSEL and cell temperatures, were obtained by sweeping the VCSEL current and measuring the amount of light intensity absorbed by the Rb in the cell. The emitted laser wavelength has a temperature and current dependence according to:

$$\lambda = \lambda_0 + \alpha \cdot (I - I_0) + \beta \cdot (T - T_0), \tag{1}$$

where λ_0 =795nm, α =0.3nm/mA, β =0.055nm/°C, I_0 =4.4mA and T_0 =25°C. The parameters α and β describe how much the laser wavelength deviates from λ_0 due to changes in the VCSEL current or temperature.

As the resonance peaks occur when the VCSEL's wavelength equals the energy level of the $^{87}Rb\ D_1$ transition ($\lambda = 795 nm$), the VCSEL's temperature can be obtained by sweeping its current and using Equation (1). The sweep period was 5 seconds, and therefore in the presence of an absorption peak, the VCSEL temperature can be indirectly measured every 5 seconds.

In addition to the VCSEL temperature the cell temperature can also be obtained by observing the absorption peak linewidth. If light with intensity J_0 is transmitted through a cell with path length L, the output light intensity can be approximated by [7]:

$$J = J_0 \cdot e^{-N\sigma L}, \tag{2}$$

where N is the number density of the alkali vapor and σ is

the absorption cross-section, which has a Lorentzian frequency distribution according to:

$$\sigma = c \cdot f_{osc} \cdot r_e \cdot \frac{\Gamma/2}{(f_0 - f)^2 + (\Gamma/2)^2}, \qquad (3)$$

where c is the speed of light, f_{osc} =0.342 is the oscillator strength for the D_1 transition of 87 Rb, r_e is the electron radius, Γ is the linewidth of the broadening, f_0 is the absorption frequency and f is the laser light frequency. The rubidium number density can be written as N= P/(k_B ·T), where k_B is the Boltzmann constant, T is the Rb vapor temperature and P is the Rb vapor pressure. The Rb vapor pressure can be obtained in Torr from [8]:

$$\log_{10} P_{Rb} = 15.88253 - \frac{4529.635}{T} + 0.00058663 \cdot T - 2.99138 \cdot \log_{10} T$$
 (4)

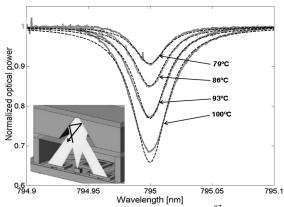


Figure 3: Normalized wavelength spectrum of 87 Rb D_1 transition at different temperatures (solid) compared to pressure broadened absorption model (dashed).

Fig. 3 shows a comparison between the described pressure broadened absorption model and the normalized ⁸⁷Rb D₁ line absorption profile measured at different temperatures. The insert in Fig. 3 shows the laser pathway inside the cell [9] (L=2.15mm).

To be able to accurately reproduce the experimental conditions in the FE model, the device was first operated in a steady-state and the initial temperatures were recorded. All measurements were taken at a sampling frequency of 100Hz with the TEC operating at a constant current of 2A, yielding a TEC temperature of -30°C.

Two different sets of measurements were collected under slightly different conditions. In the first case, measurements were taken when the heater cooled down from 195°C to 185°C. In the second set of experiments, the heater temperature was increased from 180°C to 205°C.

Measurements Filtering

The filter used in the direct experimental measurements is described in [10] and is widely used in experiments involving multi-exponential decays with time-constants spanning several orders of magnitude (for example, MEMS thermal responses). In such cases, uniform

sampling of thermal transients delivers an oversampling of the steady state. To solve this problem and reduce the noise level, a logarithmic time-scale transformation is applied to these transients. In the experiments under consideration, the filtered transients are chosen to have 40 points per decade resulting in a total of 191 and 197 samples for the first and second set of measurements, respectively.

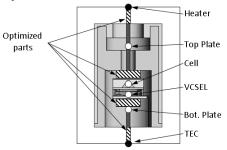


Figure 4: Cross section of NMR Gyroscope showing controlled nodes for deriving the thermal model and optimized parts.

*Table 1: Material Properties**.

Material	Thermal Conductivity	Specific Heat Capacity	Density
	[W/(m·K)]	$[J/(kg \cdot K)]$	$[kg/m^3]$
Silicon	163	703	2330
Air	0.026	1100	11.7
Polyimide HD4110	0.2	1100	1300
Loctite 1C-LV	2	1500	1300
Pyrex 7440	1.2	730	2200
Copper	400	385	8700
Amethyst SL	0.5	1100	1100

*Air has temperature dependent properties, shown values correspond to 25°C.

THERMAL MODEL

An FE model that reproduces the real prototype geometry was developed using COMSOL®. In order to reduce the complexity of the FE model, the external heater, the TEC, and the internal temperature sensors were omitted from the model. The heating and cooling copper wires were limited to 2mm in length, as shown in Fig. 4. The heater and TEC temperatures obtained with the experimental setup were applied as loads in the FE model (black dots in Fig. 4). The material properties that were used are shown in Table 1.

Optimization

Since a simplified FE model was used the thermal conductivities of some parts of the device needed to be optimized in order to match the simulation results with the measurements. The four parts chosen are shown in Fig. 4, including the top portion of the heating wire, both copper plates, and the bottom portion of the cooling wire. The optimization process [11] gives us the four thermal conductivities (see Table 2) which minimize the steady-

state temperature errors between the FE model and measurements (see Table 3).

Table 2: Optimized Thermal Conductivities

Location	Thermal Conductivity [W/(m·K)]
Top Portion of heating wire	40.4
Top Copper Plate	5.5
Bottom Copper Plate	401
Bottom Portion of cooling wire	21.8

Table 3: Steady State Optimization Results.

	Set 1			Set 2		
	Simul.	Meas.	Error	Simul.	Meas.	Error
Top Plate	118.8	117.8	1.0	130.8	130.9	-0.1
Cell	91.0	92.0	-1.0	100.9	98	2.9
VCSEL	53.2	53.0	0.2	60.6	59.5	1.1
Bot. Plate	50.1	53.7	-3.6	57.3	59.6	-2.3

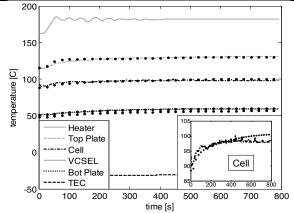


Figure 5: Illustration of matching between direct experimental measurements (lines) and optimized FE model (dotted lines) at chosen locations of the NMR Gyroscope.

Table 4: Errors comparing FE model and measurements.

Location	RMS error		ABS error		Steady-State error [°C]	
	set 1	set 2	set 1	set 2	set 1	set 2
Top Plate	1.3	1.3	1.6	4.3	1.2	0.4
Cell	0.7	1.5	1.7	5.0	0.8	2.4
VCSEL	0.5	0.5	1.4	1.2	0.5	0.8
Bot. Plate	3.5	3.2	4.2	3.8	3.3	2.7

A comparison between the thermal transients of the FE model and the experimental results for the second measurement set is shown in Fig. 5, with a close-up view of the cell in the inset picture. A summary of the errors for both measurement sets is given in Table 4. Note that the heater and TEC are not included in Table 4, since their errors are zero (were used as inputs in the FE model).

COMPACT MODEL

Once the FE model has been optimized to match the experimental measurements, the next step is to build a

CTM from the FE model. A CTM is a reduced order model which replicates with high accuracy the thermal behavior of the FE model, and in turn the physical device. The methodology is based on a Sequential Forward Selection (SFS) process described in [12]. The CTM is represented as an RC network using a thermo-electrical analogy. For the presented device, the resulting CTM has a total of 12 nodes.

Table 5 shows a summary of RMS, absolute, and steady-state errors when comparing the FE model step response transients to the obtained CTM results. The large absolute errors correspond to the central portion of the step response transients. The steady-state errors are below 0.1°C and the RMS errors are approximately 0.15°C for all four transients.

Table 5: Errors comparing CTM and FE model.

Location	RMS error	ABS error	Steady-State error
	[°C]	[°C]	[°C]
Top Plate	0.15	0.62	-0.008
Cell	0.17	0.75	0.006
VCSEL	0.14	0.39	0.021
Bot. Plate	0.13	0.36	0.051

DISCUSSION

The ultimate step of the presented process is comparison of the CTM with the experimental measurements (Table 6). This comparison highlights that the largest contribution to the CTM error is the FE model optimization step. The FE model currently does not match the experimental measurements with sufficient accuracy. Better control of the boundary conditions by utilizing an improved experimental setup needs to be achieved to reduce the uncertainties of the FE model. In addition, an accurate FE model that reproduces more details of the physical device can be implemented to reduce these errors.

Table 6: Errors comparing CTM model and measurements.

Location	RMS error		ABS error		Steady-State error [°C]	
	set 1	set 2	set 1	set 2	set 1	set 2
Top Plate	1.0	1.0	1.6	4.2	1.2	0.4
Cell	0.6	1.7	2.0	5.0	0.7	2.3
VCSEL	0.8	0.6	2.0	1.3	0.9	0.6
Bot. Plate	2.6	3.8	3.8	4.6	3.0	2.9

The presented results demonstrate that the methodology developed in this work can be used to indirectly obtain temperature estimates at critical locations of the device where direct measurements are not possible. Accurate predictive models are needed for a broad class of atomic MEMS, where temperature at unobservable locations of the system needs to be controlled with high accuracy (<1mK).

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