## Variational Weakly Supervised Gaussian Processes

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We introduce the first model to perform weakly supervised learning with Gaussian processes (GPs) on up to millions of instances. The key ingredient to achieve this scalability is to replace the standard assumption of MIL that the bag-level prediction is the maximum of instancelevel estimates with the accumulated evidence of instances within a bag. Given data set of *N* instances **X** = [ $\mathbf{x}_1 \dots \mathbf{x}_N$ ] = { $\mathbf{X}_1 \cup \mathbf{X}_1 \cup \dots \cup \mathbf{X}_B$ } composed of *B* disjoint partitions, called bags, and supervised by bag labels  $\mathbf{T} = [t_1, \dots, t_B]$ , we propose the following model to infer a Bayesian weakly supervised predictor

$$p(\mathbf{u}|\mathbf{Z}) = \mathcal{N}(\mathbf{u}|\mathbf{0}, \mathbf{K}_{\mathbf{Z}\mathbf{Z}}), \tag{1}$$

$$p(\mathbf{f}|\mathbf{u}, \mathbf{X}, \mathbf{Z}) = \mathcal{N}(\mathbf{f}|\mathbf{K}_{\mathbf{X}\mathbf{Z}}\mathbf{K}_{\mathbf{Z}\mathbf{Z}}^{-1}\mathbf{u},$$
(2)

$$diag(\mathbf{K}_{\mathbf{X}\mathbf{X}} - \mathbf{K}_{\mathbf{X}\mathbf{Z}}\mathbf{K}_{\mathbf{Z}\mathbf{Z}}^{-1}\mathbf{K}_{\mathbf{Z}\mathbf{X}})),$$

$$p(\mathbf{T}|\mathbf{f}) = \prod_{b=1}^{B} \text{Bernoulli}(T_b | \boldsymbol{\sigma}(\mathbf{f}_b^T \mathbf{1})), \quad (3)$$

where  $\mathbf{Z} = [\mathbf{z}_1; \dots; \mathbf{z}_P]$  with  $P \ll N$  is a tiny pseudo data set called the *inducing point set* and **u** the corresponding outputs. We refer to this model as the *Variational Weakly Supervised Gaussian Process* (*VWSGP*)<sup>1</sup>. Here, Equations 1 and 2 constitute the sparse GP prior and Equation 3 is a Bernoulli likelihood ensuring the model to predict binary outputs. Thanks to the sum term  $\mathbf{f}_b^T \mathbf{1}$  in the likelihood, this model can be trained by closed-form variational inference updates. Hence, keeping all parameters but one fixed, the remaining parameter can be analytically fit to the global optimum. This virtue leads to charmingly fast convergence, fitting perfectly to large-scale learning setups.

We evaluate our VWSGP on the Pascal VOC '07 benchmark and two medical image analysis applications: i) Diabetic Retinopathy screening (DR), and ii) metastatic tumor detection from histopathology images of lymph node tissues (Lymph). While VOC '07 consists of 19M instances (2000 region proposals per image), DR and Lymph have 361K and 1M instances, respectively. The results are summarized in Table 1. Our model proves to outperform various scalable MIL algorithms, as well as state-of-theart adaptations of deep learning to weakly supervised learning.

Table 1:	Bag-level	average	precision	scores of	n
two medi	ical data se	ets.			

	VOC'07	DR	Lymph
VWSGP (Ours)	83.7	0.98	0.68
VGG-S [1]	82.4	-	-
<b>DMIL</b> [4]	75.5	-	-
<b>mi-FV</b> [3]	-	0.92	0.48
<b>e-MIL</b> [2]	-	0.93	0.61

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<sup>&</sup>lt;sup>1</sup>The source code of our model is publicly available under https://github.com/melihkandemir/vwsgp