A Surrogate-Assisted and Informed Linkage Aware GA

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ABSTRACT

We present a genetic algorithm that combines linkage learning, surrogate models and informed operators. Linkage learning aims at measuring and exploiting interdependence of groups of genes. Surrogate models are fitness approximators to ease the task of calculating true fitness values. Informed operators generate, evaluate and rank a set of solutions according to their fitness model to return their most fit solution. Our described approach provides on-line perturbation based linkage learning and informed linkage exploitation with novel, specialized operators. Results of experimental runs on several synthetic fitness function compositions are provided to demonstrate significant improvement of the final result quality compared to a conventional GA setup.

Categories and Subject Descriptors

I.2.6 [ARTIFICIAL INTELLIGENCE]: Learning— Parameter learning; I.2.8 [ARTIFICIAL INTELLIGENCE]: Problem Solving, Control Methods, and Search— Heuristic methods

Keywords

Genetic Algorithms, Surrogate Models, Linkage, Operators

1. INTRODUCTION

The type and strength of interactions of problem variables in the context of global optimization is referred to as linkage or epistasis, and its character and possible exploitation has been subject to continuous studies in the area of evolutionary computation (EC) [1]. Another approach to save evaluation time and enhance the quality of final individuals is surrogate-assisted evolution (SAE). Here, surrogate models (for instance neural networks [3]) work as function approximators. Informed operators, introduced in [8], work by applying a surrogate model during offspring creation to rank an offspring pool. Surrogate models have been previously employed in linkage learning [5], but only in an offline linkage learning fashion without any operators to take advantage of it. We present a novel combination of these three approaches, the surrogate-assisted and informed linkage exploiting genetic algorithm (SAILEGA), which allows for on-line perturbation based linkage learning and informed exploitation of it.

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2. LINKAGE LEARNING

One of the major goals of linkage learning is the detection of the underlying fitness function's decompositability. Formally, the fitness function F(x) in Equation 1

$$F(x) = \sum_{i=0}^{k} f_i(x_{s_i})$$
(1)

can be additively decomposed into k sub-functions, where the whole chromosome x is split into subchromosomes x_{s_i} , and where s_i defines which genes belong to the corresponding subchromosome (with permitted overlapping). Such a set of genes s_i is called a linkage group. SAILEGA¹ implements surrogate-assisted and regular versions of LINC-R [9] and LIMD [4]. To mend linkage estimation errors, we define existing linkage for two genes only if both surrogate-assisted LINC-R and LIMD return a successful linkage observation.

3. SURROGATE MODELS

All encountered (unique) individuals are stored in a population storage, which is clustered with X-means [6] every generation. A SVM for regression using Weka [2] with the Pearson VII function-based universal kernel (PUK) is created for each cluster that is represented by at least one member of the current population. Every generation, one member of each current cluster is subject to linkage analysis.

4. CROSSOVER OPERATORS

SAILEGA incorporates four novel crossover operators combining the merits of linkage exploitation, SAE and informed operators with the aforementioned SVMs and a crossover selector following GADO's [7] guideline. The informed linkage group line (ILGL) crossover is based on the double line crossover [7], it is especially helpful in regions of ellipsoidal space, the algorithm is given in Figure 1. The informed guided linkage group (IGLG) crossover is based on the guided crossover [7]. Similarly as with the ILGL crossover, each linkage group receives its own ratio variable r to let linked genes exploit the search space in the same direction. The informed linkage inter group (ILIG) crossover takes the idea of the uniform crossover to the level of linkage groups to keep linked genes from being disrupted while enabling mixing of genetic material. The informed linkage inter-intra group (ILIIG) crossover performs ILIG's recombination and then shuffles the offspring alleles of each linkage group.

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 $^{^{1}\}mathrm{Our}$ GA is built on top of the Watchmaker Framework 0.7.1: http://watchmaker.uncommons.org/

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1: p1, p2 \leftarrow parentSelection(currentPopulation)

2: offspring \leftarrow empty offspring

3: allGroups \leftarrow set of linkage groups \cup nonlinked group

4: for all group \in allGroups do

5: r \leftarrow getRandom(-2, 2)

6: for all gene \in group do

7: newGene = r * p1_{gene} + (1 - r) * p2_{gene}

8: offspring_{gene} \leftarrow newGene

9: end for

10: end for
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Figure 1: The ILGL crossover algorithm.

Table 1: Fitness Functions Function Definition					
F_1 : Sphere ₄	$\sum_{i=1}^{D} (x_i + 4)^2$				
F_2 : Sphere ₋₄	$\sum_{i=1}^{D} (x_i - 4)^2$				
F_3 : Rosenbrock	$\sum_{i=2}^{i-1} (100(x_1 - x_i^2)^2 + (x_i - 1)^2)$				
F_4 : Schwefel 1.2	$\sum_{i=1}^{i-1} (\sum_{i=1}^{i} x_j)^2$				
F_5 : Rastrigin	$10D + \sum_{i=1}^{D} (x_i^2 - 10\cos(2\pi x_i))$				

5. EXPERIMENTAL SETUP

Three configurations C are run 50 times on each composite function as given in Tables 1 and 2. $C_{default}$ uses surrogate models, C_{noSu} only calls the true fitness function for any evaluations and C_{1point} only applies the 1-point crossover. Their remaining setup is identical to the GA in [5].

6. EXPERIMENTAL RESULTS

Table 2 shows the mean of the final individuals and the corresponding standard deviation as well as one-tailed t-test results to compare $C_{default}$ and C_{noSu} against C_{1point} . On all runs, both $C_{default}$ and C_{noSu} clearly outperform C_{1point} , showing that their configuration leads the GA to vastly superior solutions. Furthermore, with expensive fitness functions, C_{noSu} may take much more time than $C_{default}$ due to C_{noSu} 's solely calls to the actual fitness function.

Table 2: Final Minimization Results

Config	Function ^{Dim}	Mean	Std. Dev.	p-value
$C_{default}$	$F_1^7 + F_2^7$	0.0027	7.68e-4	1.168e-10
C_{noSu}	$F_1^7 + F_2^7$	0.002	0.00036	1.15e-10
C_{1point}	$F_1^7 + F_2^7$	1.32	1.17	-
$C_{default}$	$F_3^6 + F_2^8$	4.06	3.75	2.911e-10
C_{noSu}	$F_3^6 + F_2^8$	1.94	2.27	7.383e-12
C_{1point}	$F_3^6 + F_2^8$	20.44	14.99	-
$C_{default}$	$F_4^4 + F_5^4 + F_2^6$	2.03	1.43	2.567e-10
C_{noSu}	$F_4^4 + F_5^4 + F_2^6$	0.08	0.2	< 2.2e-16
C_{1point}	$F_4^4 + F_5^4 + F_2^6$	4.86	2.42	-

7. CONCLUSION

In this paper, we provide a combination of surrogate models, linkage learning and informed linkage exploiting operators. The presented method SAILEGA is composed of an on-line perturbation based linkage learning module, an application of X-means clustering to dynamically adjust its surrogate model set of SVMs for regression, and an arsenal of novel informed linkage exploiting recombination operators. These operators are designed in a modular way such that it is possible to use them in a non-surrogate setting or without a pool of created offspring, making them potentially useful for an implementation by other optimizers. We empirically demonstrated that SAILEGA can successfully exploit linkage of genes from fitness functions to significantly improve final result quality.

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