

The Gradient Flow Coupling in Minimal Walking Technicolor

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We present a measurement of the running coupling in $SU(2)$ with two adjoint fermions in the Yang-Mills gradient flow scheme. The simulations are performed with Schrödinger Functional boundary conditions using an improved HEX-smearred Wilson fermion action. We obtain a step scaling function by defining the coupling at a scale relative to the finite size of the lattice. We find a continuum limit with a non-trivial infrared fixed point.

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[†]A footnote may follow.

1. Introduction

In the space of gauge field theories coupled to fermions, there is a range in the number of colors and fermion flavors where the β -function has a non-trivial zero in the infrared. These conformal theories have applications in model building beyond the standard model, particularly in technicolor theories, where the electroweak symmetry is broken by a spontaneous symmetry breaking in a strongly interacting model. They are also interesting from the theoretical point of view of understanding the structure of gauge field theories.

When the number of fermion flavors is large, close to the upper limit of the conformal window, the coupling at the fixed point is small and perturbation theory is expected to apply. When approaching the lower limit, however, the fixed point coupling grows and non-perturbative methods are needed. The β -function can be studied in lattice field theory by studying the behavior of a renormalized coupling when the renormalization scale is changed [1].

Close to the fixed point the coupling runs very slowly and high accuracy is required to discern the direction of the running and to find the fixed point. To improve the accuracy, we study the gradient flow method for measuring the coupling. The method has been studied previously in several models and with varying boundary conditions [2–5]. Compared to the more standard Schrödinger Functional method, the statistical accuracy of the gradient flow method is higher. However, in the form presented here, the method also has larger discretization errors.

We study the coupling in the SU(2) gauge theory coupled with two fermions in the adjoint representation. In previous studies using the Schrödinger functional method the model has exhibited a non-trivial infrared fixed point at a relatively small coupling [6–18]. It is also relatively fast to simulate due to its minimal gauge and fermion content, making it a good test case for the gradient flow method.

2. Gradient Flow Coupling

Several applications of the gradient flow in lattice simulations have been studied recently [19–24]. In gradient flow, the gauge field is allowed to develop in a fictitious flow time along the gradient of an action. The flow defined by the continuum gauge action is given by

$$\begin{aligned}\partial_t B_{t,\mu} &= D_{t,\mu} B_{t,\mu\nu}, \\ B_{0,\mu} &= A_\mu \\ B_{t,\mu\nu} &= \partial_\mu B_{t,\nu} - \partial_\nu B_{t,\mu} + [B_{t,\mu}, B_{t,\nu}].\end{aligned}$$

Here $B_{t,\mu}$ is the flow field parametrized by the flow time t , and A_μ is the original gauge field. The flow naturally smooths the field, moving toward the minimum of the action. Remarkably, the flow field $B_{t,\mu}$, at any finite flow time, is a finite renormalized field. The correlators of the flow field are automatically renormalized and therefore encode physical properties of the theory [25].

As a representative case, the observable

$$\langle E(t) \rangle = \frac{1}{4} \langle G_{\mu\nu}(t) G_{\mu\nu}(t) \rangle$$

was studied in [20]. To the leading order in perturbation theory, it has the form

$$\langle E(t) \rangle = Ng^2/t^2 + \mathcal{O}(g^4).$$

The observable can therefore be used to define a renormalized coupling, [2]

$$g_{GF}^2 = \frac{t^2 \langle E(t) \rangle}{N}.$$

We study the running of the coupling with respect to the renormalization scale by measuring the coupling with several physical lattice sizes. To quantify the change we use the step scaling function [26]

$$\begin{aligned} \Sigma(u, a/L) &= g_{GF}^2(g_0, 2L/a) \Big|_{g_{GF}^2(g_0, L/a)=u} \\ \sigma(u) &= \lim_{a \rightarrow 0} \Sigma(u, a/L). \end{aligned} \quad (2.1)$$

The step scaling function describes how the coupling evolves when the linear size of the system changes from L to $2L$. The gradient flow introduces another length scale, $l = \sqrt{8t}$, which we also need to change when varying the lattice size. In practice we choose $l = 0.5L$.

We use the same highly improved action as in our Schrödinger functional calculations to allow direct comparison of the results. Preliminary results from this study have already been published [18] and the full results will be published soon. The action is given by

$$S = (1 - c_g)S_G(U) + c_g S_G(V) + S_F(V) + c_{SW} \delta S_{SW}(V),$$

where $S_G(U)$ is the standard Wilson gauge action and $S_G(V)$ is the gauge action calculated using HEX smeared links [27]. The gauge smearing parameter c_g is set to 0.5. $S_F(V)$ and $\delta S_{SW}(V)$ are the Wilson fermion action and the clover correction using HEX smeared links. The clover coefficient takes the tree level value $c_{SW} = 1$.

We perform the simulations using Schrödinger functional boundary conditions. They have some advantages over fully periodic boundary conditions: The gauge field has a unique global minimum, which simplifies perturbation theory and removes non-even orders of coupling from the perturbative expansion of the β -function [3]. The boundary conditions also alleviate problems with massless fermions. For the gauge fields they are given by

$$\begin{aligned} U_k(x) &= 1, \text{ when } x_0 = 0, L \\ U_\mu(x + L\hat{k}) &= U_\mu(x). \end{aligned}$$

The fermion fields are set to zero at the time boundaries and have twisted periodic boundary condition in the spatial directions:

$$\psi(x + L\hat{k}) = e^{i\pi/5} \psi(x).$$

Since the translational symmetry is now broken in the time direction, we avoid any unnecessary discretization errors by only using the middle time slice for the measurement,

$$g_{GF}^2 = \frac{t^2 \langle E(t, x_0) \rangle}{N}, \quad x_0 = L/2, \quad t = c^2 L^2 / 8. \quad (2.2)$$

Because of the improved lattice action, the normalization factor N is unknown on the lattice. We use the continuum value $N = 288.527$.

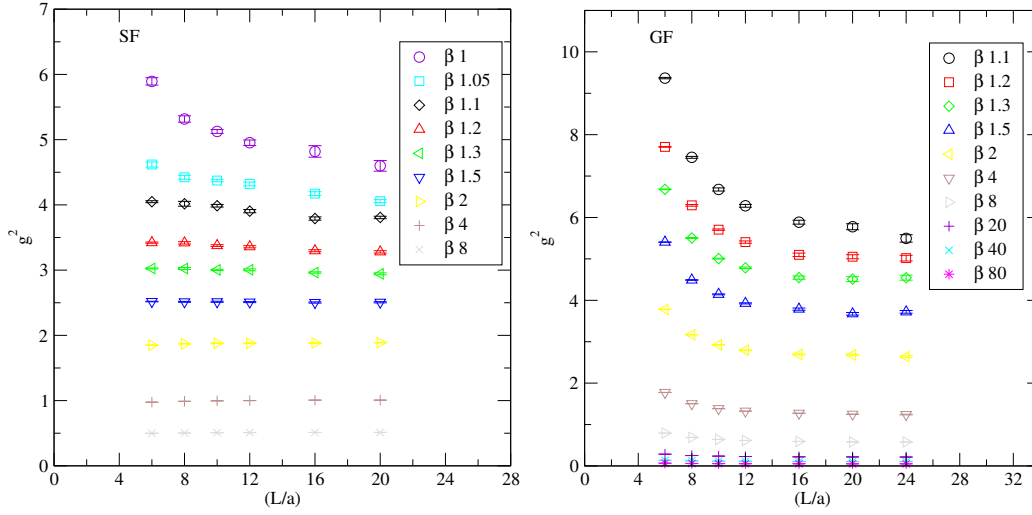


Figure 1: The measured values of the coupling using the Schrödinger functional and the gradient flow methods. The plot on the left shows the SF coupling and the plot on the right shows the GF coupling.

	SF		GF		
L/a	20	20	20	20	24
$\beta = 4/g_0^2$	2	1.2	2	1.2	1.2
N_{traj}	151203	158828	14065	13693	9648
$(\delta g^2/g^2)_{\text{statistical}}$	0.0059	0.0109	0.0112	0.0121	0.0150
$(\delta g^2/g^2)^2 N_{\text{traj}}$	5.225	18.75	1.765	2.048	2.250
t_{traj}	1221.11	1277.04	585.58	769.64	1787.13

Table 1: The statistical errors of the SF and the GF couplings at two values of β with large lattice sizes. $(\delta g^2/g^2)^2 N_{\text{traj}}$ estimates the number of trajectories required to reach a given statistical error.

L/a	6	8	10	12	16	20	24
SF, $\beta = 2$	0.093	0.259	0.265	0.880	3.791	5.225	
GF, $\beta = 1.2$	0.577	0.373	0.988	0.885	1.191	2.048	2.250

Table 2: The volume dependence of the error estimate $(\delta g^2/g^2)^2 N_{\text{traj}}$ in the SF method and the GF method.

3. Results

We compare the gradient flow coupling with the Schrödinger functional coupling measured in the same model. The Schrödinger functional and gradient flow couplings are measured with constant values of the bare coupling. All bare parameters, except the boundary conditions, are identical in the two sets of simulations.

It is clear already from the lattice values given in figure 1, that the gradient flow coupling has large discretization errors. The coupling should depend only little on the scale, especially close to the fixed point, but the gradient flow coupling shows strong dependence on the lattice size. Naturally the figure does not tell us how the discretization errors approach the continuum limit.

We also see that the statistical errors are roughly equal. The SF measurements are calculated

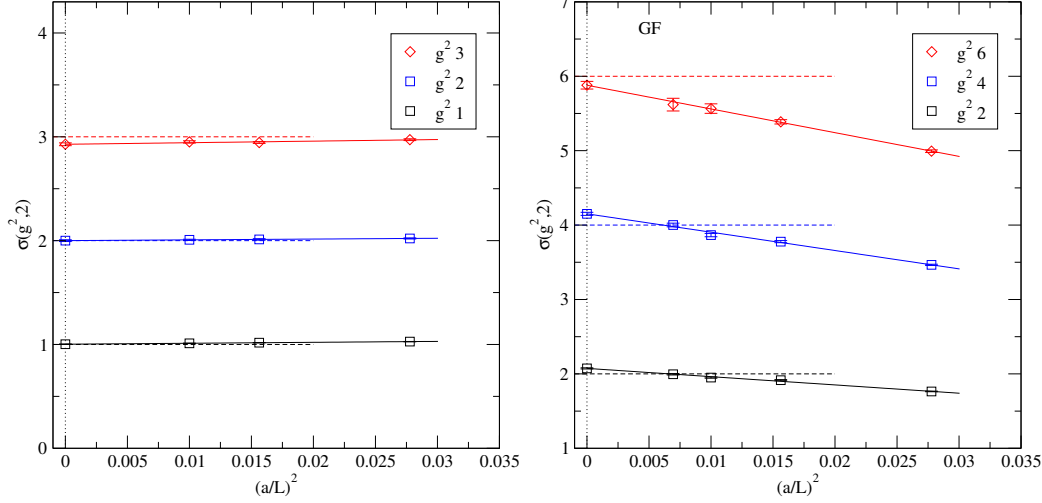


Figure 2: The continuum extrapolation of the step scaling function. The left hand side shows the SF case and the right hand side shows the GF case.

from around 150 000 HMC trajectories of length 2 and the GF measurements from around 15 000 trajectories of length 1. The behavior of statistical errors is shown in table 1 for two values of β and L/a . It is clear that the GF coupling requires less statistics for the same accuracy. The statistical accuracy of the GF coupling also scales better with the lattice size, as table 2 demonstrates.

For a more robust comparison we should take a continuum limit of the measurements. We achieve this by calculating the step scaling function $\Sigma(u, a/L)$ in equation 2.1 and taking a continuum limit keeping the renormalized coupling squared, u , constant. Since the measurements are calculated at constant bare coupling, we use an interpolating function to find the value at a given renormalized coupling. For the SF and GF couplings respectively, the interpolating functions are given by

$$g_{SF}^2(g_0) = g_0^2 \frac{\prod_{k=1}^3 (1 + a_k g_0^2)}{\prod_{k=1}^3 (1 + b_k g_0^2)}, \quad g_{GF}^2(g_0) = a_0 g_0^2 \frac{\prod_{k=1}^2 (1 + a_k g_0^2)}{\prod_{k=1}^3 (1 + b_k g_0^2)}. \quad (3.1)$$

Note that at first order, $g_{GF}^2(g_0, L/a) = a_0(L/a)g_0^2$. Thus, in the continuum limit, $a_0 = 1$, and any deviation represents a discretization error that would have been removed if we had used a lattice result for the normalization constant N in equation 2.2. We can therefore define a scaled version of the GF coupling by dividing by $a_0(L/a)$.

Using the interpolating functions we can now calculate $\Sigma(u, a/L)$ at a constant u . We estimate the continuum limit using a second order extrapolation, $\Sigma(u, a/L) = \sigma(u) + c(a/L)^2$. We show the continuum extrapolation for three representative values of u in figure 2 and the continuum value $\sigma(u)$ in figure 3.

The continuum extrapolation is steeper in the GF measurement, but nevertheless consistent with the second order extrapolation. Both measurements show a fixed point, but the GF method gives a larger coupling at the fixed point. This is partially due to the difference in the two schemes, but also suggests that there may be higher order discretization effects influencing the results.

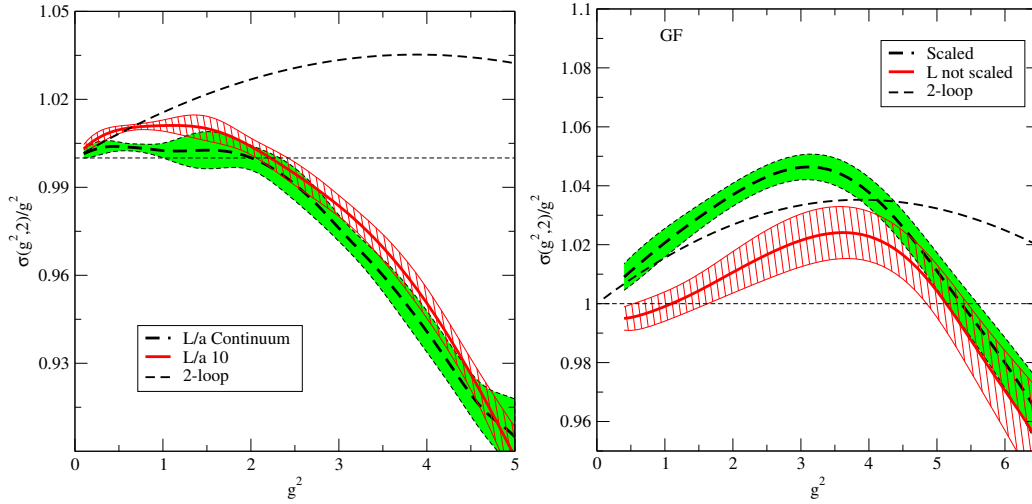


Figure 3: The scaled continuum step scaling function $\sigma(g^2, 2)/g^2$ using the SF and GF methods. On the left: The black line with the green error band corresponds to the continuum result. The red line with the hashed band correspond to the lattice result at the largest lattice size $L/a = 10$. The black dashed line gives the 2-loop perturbative result. On the right: The black line with the green band correspond the continuum limit with the coupling scaled with $a_0(L/a)$ at each lattice size. The red line with the hashed error band corresponds to the continuum result without scaling. The black dashed line gives the 2-loop perturbative result.

4. Conclusions

We have studied the gradient flow coupling in the SU2 gauge field theory with two fermions in the adjoint representation with Schrödinger functional boundary conditions. We find that the measurement is in qualitative agreement with the previous studies using the Schrödinger functional method. The results are promising, as the statistical errors are much smaller than in the Schrödinger functional coupling and scale better with the lattice size. The increased discretization effects present a problem, which may be alleviated using improved flow equations or definitions of the measurable $\langle E(t) \rangle$. It is also worth studying the effects of the boundary conditions on the coupling.

References

- [1] M. Luscher, P. Weisz and U. Wolff, Nucl. Phys. B **359** (1991) 221.
- [2] Z. Fodor, K. Holland, J. Kuti, D. Negradi and C. H. Wong, PoS LATTICE **2012** (2012) 050 [arXiv:1211.3247 [hep-lat]].
- [3] P. Fritsch and A. Ramos, arXiv:1301.4388 [hep-lat].
- [4] P. Fritsch and A. Ramos, arXiv:1308.4559 [hep-lat].
- [5] A. Ramos, arXiv:1308.4558 [hep-lat].
- [6] S. Catterall and F. Sannino, Phys. Rev. D **76**, 034504 (2007) [arXiv:0705.1664 [hep-lat]].
- [7] A. J. Hietanen, J. Rantaharju, K. Rummukainen and K. Tuominen, JHEP **0905**, 025 (2009) [arXiv:0812.1467 [hep-lat]].

- [8] L. Del Debbio, A. Patella and C. Pica, Phys. Rev. D **81**, 094503 (2010) [arXiv:0805.2058 [hep-lat]].
- [9] S. Catterall, J. Giedt, F. Sannino and J. Schneible, JHEP **0811** (2008) 009 [arXiv:0807.0792 [hep-lat]].
- [10] A. J. Hietanen, K. Rummukainen and K. Tuominen, Phys. Rev. D **80**, 094504 (2009) [arXiv:0904.0864 [hep-lat]].
- [11] F. Bursa, L. Del Debbio, L. Keegan, C. Pica and T. Pickup, Phys. Rev. D **81**, 014505 (2010) [arXiv:0910.4535 [hep-ph]].
- [12] L. Del Debbio, B. Lucini, A. Patella, C. Pica and A. Rago, Phys. Rev. D **80**, 074507 (2009) [arXiv:0907.3896 [hep-lat]].
- [13] L. Del Debbio, B. Lucini, A. Patella, C. Pica and A. Rago, Phys. Rev. D **82**, 014510 (2010) [arXiv:1004.3206 [hep-lat]].
- [14] L. Del Debbio, B. Lucini, A. Patella, C. Pica and A. Rago, Phys. Rev. D **82**, 014509 (2010) [arXiv:1004.3197 [hep-lat]].
- [15] F. Bursa, L. Del Debbio, D. Henty, E. Kerrane, B. Lucini, A. Patella, C. Pica and T. Pickup *et al.*, Phys. Rev. D **84**, 034506 (2011) [arXiv:1104.4301 [hep-lat]].
- [16] T. DeGrand, Y. Shamir and B. Svetitsky, Phys. Rev. D **83**, 074507 (2011) [arXiv:1102.2843 [hep-lat]].
- [17] J. Giedt and E. Weinberg, Phys. Rev. D **85**, 097503 (2012) [arXiv:1201.6262 [hep-lat]].
- [18] J. Rantaharju, K. Rummukainen and K. Tuominen, arXiv:1301.2373 [hep-lat].
- [19] M. Luscher, Commun. Math. Phys. **293** (2010) 899 [arXiv:0907.5491 [hep-lat]].
- [20] M. Luscher, JHEP **1008** (2010) 071 [arXiv:1006.4518 [hep-lat]].
- [21] A. Deuzeman and U. Wenger, PoS LATTICE **2012** (2012) 162.
- [22] M. Luscher, JHEP **1304** (2013) 123 [arXiv:1302.5246 [hep-lat]].
- [23] H. Suzuki, PTEP **2013** (2013) 8, 083B03 [arXiv:1304.0533 [hep-lat]].
- [24] L. Del Debbio, A. Patella and A. Rago, arXiv:1306.1173 [hep-th].
- [25] M. Luscher and P. Weisz, JHEP **1102** (2011) 051 [arXiv:1101.0963 [hep-th]].
- [26] M. Luscher, R. Sommer, P. Weisz and U. Wolff, Nucl. Phys. B **413** 481 (1994) [hep-lat/9309005].
- [27] S. Capitani, S. Durr and C. Hoelbling, JHEP **0611** (2006) 028 [hep-lat/0607006].