



2 **Comment on “Transport and fate of bacteria in porous media:**
 3 **Coupled effects of chemical conditions and pore space**
 4 **geometry” by Saeed Torkzaban et al.**

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 10 effects of chemical conditions and pore space geometry” by Saeed Torkzaban et al., *Water Resour. Res.*, 45, XXXXXX,
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13 [1] The article by *Torkzaban et al.* [2008] provides
 14 interesting data supporting the expectation that a significant
 15 fraction of colloids (bacterial cells in this case) are retained
 16 in porous media without actual attachment to collector
 17 surfaces. However, the authors present a theoretical approach
 18 that warrants elaboration regarding its relationship to pre-
 19 viously existing approaches.

20 [2] The theoretical analyses provided by the authors
 21 involved a balance of the driving and resisting torques
 22 acting on an immobilized colloid to determine whether
 23 rolling (and by extension detachment) of the colloid would
 24 be initiated, as described in previous articles [e.g., *Hubbe*,
 25 1984, 1985; *Bergendahl and Grasso*, 2000; *Li et al.*, 2005].
 26 In this approach, the contact area between the colloid and
 27 the surface (its radius) provides a lever arm, which along
 28 with the adhesion force comprises the torque that resists
 29 detachment (Figure 1). A torque driving detachment is
 30 generated by fluid drag acting at a point somewhat above
 31 the centroid of the colloid ($1.399 \times$ colloid radius) [*Sharma*
 32 *et al.*, 1992]. This torque balance has been traditionally
 33 applied to colloids immobilized to surfaces, i.e., in tradi-
 34 tional parlance, where the colloid has overcome any repul-
 35 sive energy barrier, and has come into physical contact with
 36 the surface (Figure 1). For colloids that overcome the
 37 energy barrier, the adhesive torque typically dominates,
 38 and the colloid is typically considered irreversibly attached
 39 (a perfect sink boundary).

40 [3] *Torkzaban et al.* [2008] and previously *Torkzaban*
 41 *et al.* [2007] have applied this particular torque balance in
 42 the context of colloids associated with surfaces via second-
 43 ary energy minima. They approximate the adhesion force
 44 with the attractive force experienced in the secondary
 45 energy minimum interaction; whereas previous approaches
 46 have approximated the adhesion force with the stronger
 47 attractive force experienced in the primary energy minimum
 48 (Figure 2).

49 [4] The approach of *Torkzaban et al.* [2007, 2008]
 50 deserves further discussion because previously published

51 force and torque balances for secondary energy minimum 51
 52 associated colloids allow translation in response to net 52
 53 forces (fluid drag, diffusion, gravitation, van der Waals, 53
 54 and electric double layer) as well as spinning in response to 54
 55 fluid shear [*Rajagopalan and Tien*, 1976; *Prieve and Lin*, 55
 56 1980; *Yang et al.*, 1998; *Johnson et al.*, 2007]. The latter is 56
 57 built into the force balance by way of hydrodynamic 57
 58 retardation coefficients [e.g., *Rajagopalan and Tien*, 1976; 58
 59 *Johnson et al.*, 2007]. This approach is taken with the 59
 60 expectation that the energy barrier between the surface 60
 61 and the secondary energy minimum prevents physical 61
 62 contact between the colloid and stationary surface, such 62
 63 that the friction resisting colloid motion emanates from the 63
 64 fluid viscosity rather than emanating from adhesive contact 64
 65 between the colloid and the surface (Figure 3).

66 [5] The major difference in treatment of secondary 66
 67 energy minimum associated colloids in the traditional 67
 68 approaches versus that invoked by *Torkzaban et al.* [2007, 68
 69 2008] is the source of friction that resists colloid motion. In 69
 70 the traditional approach, the colloid has no adhesive contact 70
 71 with the surface, and friction emanates from the viscosity of 71
 72 the fluid in which the colloid translates and rotates; whereas 72

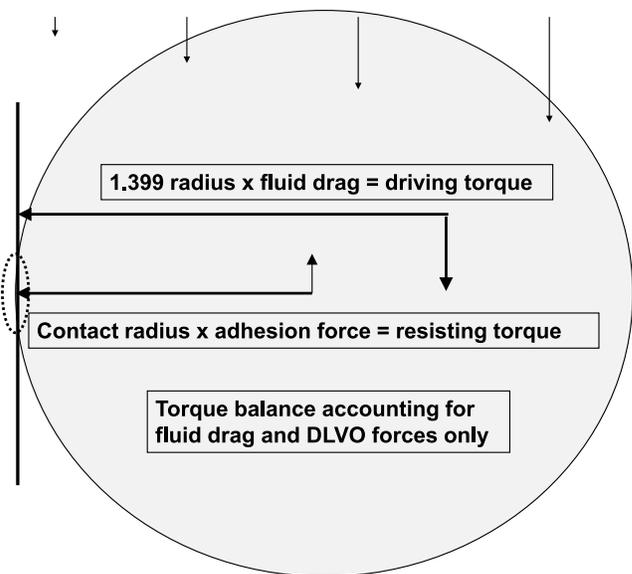


Figure 1. Schematic of colloid attached to surface with corresponding driving torque (driving detachment) originating from fluid drag (depicted by arrows at top) and resisting torque (resisting detachment) originating from adhesion.

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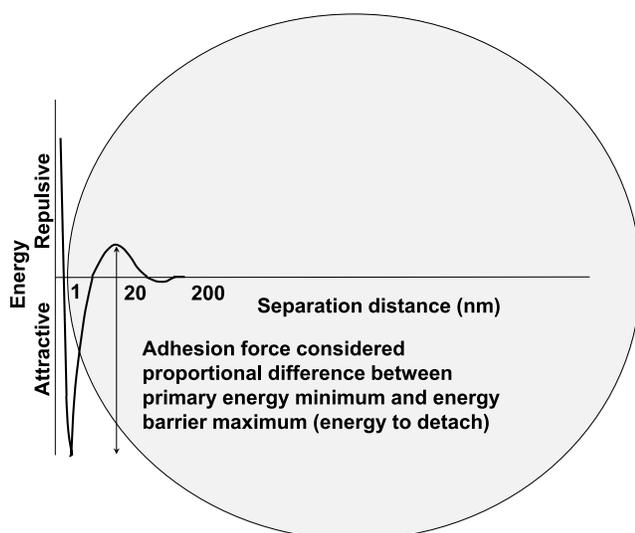


Figure 2. Schematic of colloid-surface interaction force profile superimposed on colloid of ~ 800 nm diameter associated with the surface via the primary energy minimum.

73 in the approach by *Torkzaban et al.* [2007, 2008], friction
74 emanates from adhesive contact between the colloid and the
75 surface, and colloid translation is assumed to not occur.

76 [6] A great deal of experimental data can be cited that
77 implicates retention of colloids in secondary energy minima.
78 The issue we raise is not whether retention of secondary
79 energy minimum associated colloids occurs, but rather, how
80 it occurs. Our own simulations using a traditional force and
81 torque balance for secondary energy minimum associated
82 colloids show that colloids may be retained in zones of low
83 fluid drag at the pore scale even without adhesive contact
84 with the surface [*Johnson et al.*, 2007]. In our model,
85 retention without attachment occurred in rear stagnation

zones, although these represent just one possible type of 86
zone of low fluid drag. *Torkzaban* and colleagues appear to 87
incorrectly assume that retention without attachment occurs 88
only in flow vortices. It must be noted that complete flow 89
stagnation is not necessary to produce colloid retention; 90
rather, only sufficiently low fluid drag and sufficiently high 91
secondary energy minimum attraction are required to pro- 92
duce retention, and this retention occurs without attachment 93
[*Johnson et al.*, 2007], as opposed to primary energy 94
minimum associated colloids (e.g., via heterogeneity and 95
wedging), which are genuinely attached. There are two 96
important points here: (1) retention of secondary energy 97
minimum associated colloids can be simulated in a system 98
where colloid spinning and translation are allowed, and 99
(2) spinning of the colloid in response to fluid shear does 100
not necessarily lead to reentrainment. These two points, 101
demonstrated by *Johnson et al.* [2007] for secondary energy 102
minimum associated colloids, contrast with the assumptions 103
used by *Torkzaban et al.* [2007, 2008] that (1) colloids may 104
be immobilized by secondary energy minimum interaction; 105
that is, secondary energy minimum interaction constitutes 106
adhesive contact with the surface, and (2) the initiation of 107
rolling can be equated to reentrainment. 108

[7] That physical contact is established when a colloid 109
associates with a surface via the primary energy minimum is 110
quite clear, since the equilibrium separation distance is close 111
to 0.16 nm [*Israelachvili*, 1992], where Born repulsion 112
results from the overlap of electron orbitals on the two 113
surfaces. In contrast, the secondary energy minimum 114
involves separation distances of ten to hundreds of nm, 115
and the notion of contact is much less clear. Adhesion 116
theory [e.g., *Johnson et al.*, 1971; *Derjaguin et al.*, 1975] 117
supposes physical contact between the colloid and the 118
surface, incorporating some degree of physical deformation 119
of the colloid and the surface via Young's moduli and 120
Poisson ratios to yield an adhesive contact area [see 121
Bergendahl and Grasso, 2000], such that the friction 122

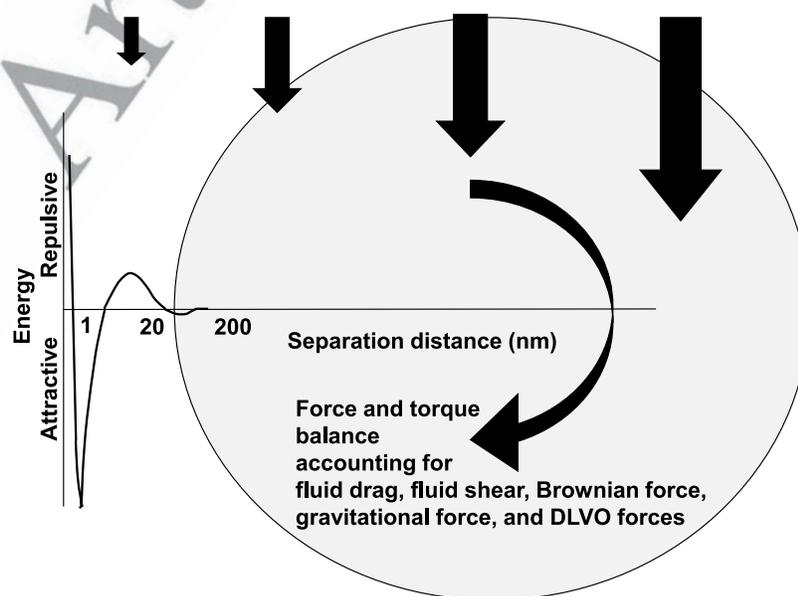


Figure 3. Schematic of colloid-surface interaction force profile superimposed on colloid of ~ 800 nm diameter associated with the surface via the secondary energy minimum. Arrows depict fluid drag force and associated torque.

123 resisting colloid motion emanates from the colloid-surface
 124 contact. The corresponding equation for contact radius
 125 provided by *Torkzaban et al.* [2007, 2008] was obtained
 126 by manipulation of equations for the contact radius and pull-
 127 off force provided by *Israelachvili* [1992, chapter 15]. That
 128 source clearly states that the equations apply to materials in
 129 adhesive contact. Although secondary energy minimum
 130 interactions (weak van der Waals) occur over definable
 131 areas on the colloid and collector surfaces [*Israelachvili*,
 132 1992], the friction resisting colloid motion in secondary
 133 energy minima has traditionally been considered to arise
 134 from the colloid-fluid interface (fluid viscosity). *Torkzaban*
 135 *et al.* [2007, 2008] depart from this traditional approach,
 136 and equate secondary energy minimum to adhesive contact,
 137 a move that warrants further discussion and exploration.
 138 This comment provides an opportunity for Torkzaban and
 139 colleagues to substantiate their stance that adhesion param-
 140 eters developed for contact are applicable to secondary
 141 energy minimum interactions.

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