Facile, rapid and efficient isolation of circulating tumor cells using aptamer-targeted magnetic nanoparticles integrated with a microfluidic device

Laleh Rafiee^{a‡}, Abolghasem Abbasi Kajani^{b‡}, Mohamadmahdi Samandari^{a,c}, Masoud Ayatollahi

Mehrgardi^d, Bahare Zarrin^a, Shaghayegh Haghjooy Javanmard^{a,*}

^a Applied Physiology Research Center, Cardiovascular Research Institute, Isfahan University of Medical Sciences,

Isfahan, 81746-73461, Iran

^b Department of Biotechnology, Faculty of Biological Science and Technology, University of Isfahan, Isfahan, 81746-73441, Iran

° Department of Biomedical Engineering, University of Connecticut, Farmington, CT 06030, USA

^d Department of Chemistry, University of Isfahan, Isfahan, 81746-73441, Iran

[‡]These authors contributed equally to this work.

* Corresponding authors:

- Shaghayegh Haghjoo Javanmard, Email: <u>sh_haghjoo@med.mui.ac.ir</u>



Fig. S1. The viability of HUVEC (yellow) and MCF-7 (blue) cells after exposure with different concentrations of MNPs.



Fig. S2. Effect of the incubation time (A) and MNPs concentration (B) on the capture efficiency of MCF-7 cells by using APT-MNPs.



Fig. S3. The optical microscope image of captured MCF-7 cancer cells by Apt-MNPs.



Fig. S4. The viability and proliferation of magnetically captured MCF-7 cancer cells. (A) Immediately after capturing and (B) 2 weeks after *in vitro* culture.



Fig. S5. The optical microscope images of magnetically captured cells from the blood samples of breast cancer patient (A) and healthy donor (B).

Table S1. The number of isolated CTCs from the peripheral blood samples of breast cancer patients using the APT-MNPs and EasySepTM Kit.

samples	APT-MNPs	EasySep™ Kit	Stage
1	2	2	4
2	2	1	4
3	2	1	4
4	1	0	4
5	2	0	4