Down-Regulation of miR-200c and Up-Regulation of miR-30c Target both Stemness and Metastasis Genes in Breast Cancer

Mahsa Rahimi, M.Sc.1,2, Ali Sharifi-Zarchi, Ph.D.2, Nosratollah Zarghami, Ph.D.1, Lobat Geranpayeh, MD.3, Marzieh Ebrahimi, Ph.D.2*, Effat Alizadeh, Ph.D.4*

1. Department of Medical Biotechnology, Faculty of Advanced Medical Sciences, Tabriz University of Medical Sciences, Tabriz, Iran
2. Department of Stem Cells and Developmental Biology, Cell Science Research Center, Royan Institute for Stem Cell Biology and Technology, ACECR, Tehran, Iran
3. Department of Surgery, Sina Hospital, Tehran University of Medical Sciences, Tehran, Iran
4. Drug Applied Research Center, Tabriz University of Medical Sciences, Tabriz, Iran

*Corresponding Addresses: P.O. Box 16635-148, Department of Stem Cells and Developmental Biology, Cell Science Research Center, Royan Institute for Stem Cell Biology and Technology, ACECR, Tehran, Iran
R.O.Box: 5166653431, Drug Applied Research Center, Tabriz University of Medical Sciences, Tabriz, Iran
Emails: mebrahimi@royaninstitute.org, alizadehe@tbzmed.ac.ir

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Abstract

Objective: microRNAs (miRNAs) play important role in progression of tumorigenesis. They can target self-renewal and epithelial-mesenchymal transition (EMT) abilities in tumor cells, especially in cancer stem cells (CSCs). The objective of this study was to implement data mining to identify important miRNAs for targeting both self-renewal and EMT. We also aimed to evaluate these factors in mammospheres as model of breast cancer stem cells (BCSCs) and metastatic tumor tissues.

Materials and Methods: In this experimental study, mammospheres were derived from MCF-7 cells and characterized for the CSCs properties. Then expression pattern of the selected miRNAs in spheroids were evaluated, using the breast tumor cells obtained from seven patients. Correlation of miRNAs with self-renewal and EMT candidate genes were assessed in mammospheres and metastatic tumors.

Results: The results showed that mammospheres represented more colonogenic and spheroid formation potential than MCF-7 cells (P<0.05). Additionally, they had enhanced migration and invasive capabilities. Our computational analyses determined that miR-200c and miR-30c could be candidates for targeting both stemness and EMT pathways. Expression level of miR-200c was reduced, while miR-30c expression level was enhanced in mammospheres, similar to the breast tumor tissues isolated from three patients with grade II/III who received neo-adjuvant treatment. Expression level of putative stem cell markers (OCT4, SOX2, c-MYC) and EMT-related genes (SNAIL1, CDH2, TWIST1/2) were also significantly increased in mammospheres and three indicated patients (P<0.05).

Conclusion: Simultaneous down-regulation and up-regulation of respectively miR-200c and miR-30c might be signature of BCSC enrichment in patients post neo-adjuvant therapy. Therefore, targeting both miR-200c and miR-30c could be useful for developing new therapeutic approaches, against BCSCs.

Keywords: Metastasis, miR-200c, miR-30c, Self-Renewal, Spheroid

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Introduction

Breast cancer is the second-most prevalent cancer between females worldwide (1) and effective treatment of breast cancer is faced to a number of hurdles including resistance to therapies, metastasis and recurrence (2). There are several evidences regarding the heterogeneity of breast cancer cell population, initiated from a very slight subset of cells named cancer stem cells (CSCs) (3). CSCs with self-renewal capacity are responsible for initiation of tumorigenesis in immunodeficient models (4) as well as maintenance and clinical outcomes of treatments (5). Although CSCs play central role from clinical points of view, molecular mechanisms and pathways involved in their survival and maintenance has not fully been identified (6). Increasing our knowledge in the field of tumor biology could consequently lead to suggestion of effective diagnostic and prognostic methods, as well as more impressive treatment for breast cancer. Moreover, recent progress has highlighted the important role of microRNAs (miRNAs) in regulating stemness and metastasis of CSCs. In this way, several miRNAs are known to be differentially expressed in CSCs or normal stem cells, part of which has been studied in targeting genes and networks involved in cancer stemness properties (7). On the other hand, the regulatory role of miRNAs has been defined in epithelial-mesenchymal transition (EMT), as an important process through tumor progression (8). Although it is known that miRNAs could contribute to tumorigenesis as tumor suppressors or oncogenes (9, 10), the role of miRNAs targeting both self-renewal and EMT pathways in breast CSCs (BCSCs) has largely been remained unknown.

Recently, with regards to the new technologies innovation, data mining and bioinformatics approaches have tremendously been developed in the field of
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Genomic analysis large-scale endeavors created useful databases. We hypothesized that identification of common miRNAs targeting stemness-EMT network will improve our understanding of CSCs in metastatic breast cancer. To date, several investigations have been performed to find deregulated miRNA expression during EMT and metastasis of breast cancer or BCSCs. In this study, our approach is systematic analysis of combined clinical and molecular data to find common miRNAs deregulated in mammospheres, as BCSCs model, and metastatic breast cancer. To reach that, we integrated candidate miRNA expression profiles with their target mRNA gene expression data obtained from the same samples. In summary, our results led to study the important role of miR-200c and miR-30c in maintenance of stemness as well as EMT process in BCSCs. Therefore, we suggested that down-regulation of miR-200c combined with increasing level of miR-30c may be a signature of BCSCs enrichment in patients post neo-adjuvant therapy. These miRNAs may have potential to extent into both diagnostic filed, as biomarker, and therapeutic approach for BCSCs in patients who are under chemotherapy.

Materials and Methods

In this experimental study, breast cancer tissues were collected between January 2017 and January 2018, upon the approval of Farmanieh Hospital and Sina Oncologic Hospital (both from Tehran, Iran) according to local authorities. All contributors signed a written informed consent form to participate in this study. All procedures performed in studies including human patient involvements were in accordance with the ethical standards that approved by Tabriz University of Medical Sciences (5/D/25333) and Royan Institute Ethical Committee (IR.ACECR.ROYAN.REC.1396.229), as well as the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Patients histopathological information, including tumor size and depth of invasion, lymph-vascular and perineural invasion, grade and clinical tumor/node/metastasis, were recorded and pathologically staged using the tumor-nodes-metastasis (TNM) staging method (11). Informed consent was obtained from all participants included in the present study in Sina and Farmanieh hospitals, Tehran, Iran.

Seven female breast cancer patients who underwent surgery at Farmanieh Hospital and Sina Oncologic Hospital were included in this research. The inclusion criteria for selection of female patients were 25 years of age and older, from all ethnicity. Breast cancer malignancy was confirmed based on histopathological examination and immunohistochemical studies of estrogen receptor (ER) and progesterone receptor (PR) expressions, performed on surgical resection tissue samples of the tumors based on the standard methods. Three samples were undergoing neo-adjuvant therapy before sampling. Normal adjacent biopsies, as negative controls, were collected from all seven patients. For sampling, surgeons removed the tumors and small part of them was cut for cultivation, which transferred to phosphate buffer saline (PBS) containing penicillin/streptomycin and the reminding part of tissues were fixed for pathological evaluation. Adjacent breast tissues or the areas around tumor sites were removed and transferred to transferring media (PBS containing penicillin/streptomycin) in the separate tube. Later on, these samples are called as normal tissues in the present study.

Literature mining and computational analysis

First we performed a systematic literature review on PubMed and COREMINE website using the following keywords: "breast cancer tissue, stem cell, self-renewal, stemness, miRNA, metastasis or EMT". The studies with incomplete data were excluded from this analysis, providing that: i. The papers are review articles or letters, ii. Studies with insufficient or inaccessible data, and iii. Studies that are not related to CSCs and homo-sapiens. We also excluded nine articles, due to limitation to access to their full texts. Moreover, miRNA expression profiles were searched with the same keywords on NCBI GEO database. In overall, we found the most frequent miRNAs targeting the stemness and metastasis genes. Then, we used miRNA target prediction tools including TargetScan (12) and miRWalk (13), to find target genes of each candidate miRNA. We only preserved the target genes with at least two-fold expression change and P<0.05, between human breast cancer versus human normal breast (the first group) and mammospheres versus MCF-7 adherent culture (the second group). Custom R scripts were used to rank miRNAs for targeting at least three stemness and two metastasis genes. Subsequently, we computed differential expression fold-changes and P values (using two-sided Student’s t test) between breast cancers vs. normal breast (as the first group) and also mammospheres vs. MCF-7 adherent culture (as the second group). Enricher (14) and GO functional enrichment analysis on KEGG 2017 pathways were used to identify pathways and biological functions that were affected by the target genes of each miRNAs.

Cell line and monolayer culture

MCF-7 is an estrogen-dependent human breast adenocarcinoma cell line that was purchased from Iranian Biological Resource Center (IBRC), Iran. The cells were cultured in Dulbecco’s Modified Eagle Medium (DMEM, Gibco, USA) supplemented with 10% heat inactivated fetal bovine serum (FBS, Invitrogen, USA), 1% non-essential amino acid (NEAA), 2 mM L-glutamine and 1% penicillin/streptomycin (all from Life Technologies, USA) at 37°C and 5% CO₂ using standard cell culture incubator.

Formation of spheroid cultures from MCF-7

The standard tissue culture plates were covered with
poly 2-hydroxyethyl methacrylate (poly-HEMA) preventing cell attachment to plate surface. Subsequently, the monolayer MCF-7 cells were enzymatically detached into single cells suspension with trypsin (Gibco, USA) and harvested. 2×10^4 single cells were seeded at low attachment plate, in serum-free DMEM medium enriched with 20 ng/ml epidermal growth factor (EGF, Rayan Institute, Iran), 20 ng/ml basic fibroblast growth factor (bFGF, Rayan Institute, Iran), 2% B27 (Gibco, USA) and 2 mM L-Glutamine (Life Technologies, USA). The media was refreshed every 48 hours and mammospheres were formed after 14 days.

**Mammosphere-forming efficiency assay**

When the spheroids reached to about 50 μm diameters, they were accumulated by gentle centrifugation at 1000 rpm for 5 minutes, and then were enzymatically separated with trypsin. About 2×10^4 cells were plated into poly-HEMA coated six-well plates in 2000 μl of serum-free DMEM medium per well. Mammosphere-forming efficiency (MFE) was calculated by dividing the number of mammospheres, which are greater than 60 μm or larger in size in the cells seeding density per well using a microscope fitted with magnitude. All experiments on each generation of mammospheres were performed in triplicates.

**Colony-forming test**

To compare colony forming capacity of the adherent cells and mammospheres, 200 cells of each group were counted and re-plated in a complete medium containing DMEM supplemented with 10% FBS, 1% NEAA, 2 mM L-glutamine and 1% penicillin/streptomycin in six-well-plates. After 10 days, cell colonies were fixed with 4% paraformaldehyde, and stained with 0.05% crystal violet (Sigma, USA). Ultimately, the round shape colonies with more than 400 μm diameter were counted using an inverted microscope (Japan Microscope brand, Japan).

**Transmembrane migration and invasion assay**

Adherent cells and mammospheres were grown up to 80% confluence. Then adherent cells were starved in serum-free medium the day before assay. The next day, the cells were dissociated into single cells with trypsin, counted and added at 1×10^6 cells/well density onto a ceramic pestle in TRizol Reagent (Invitrogen, USA). Total RNAs with the aim of small RNA retentions were extracted from the adherent cells (as control groups) and mammospheres (as experimental groups) using TRizol reagent, according to the manufacturer’s instructions. The concentration and purity of extracted RNA were determined by UV absorbance at 260 and 280 nm (260/280 nm) in spectrophotometer. The integrity of RNA samples was checked by gel electrophoresis. 2 μg total RNA was subjected to generate complementary DNA using cDNA synthesis kit (TaKaRa, Japan), according to the manufacturer’s instructions. Expression level of stemness and metastasis genes was evaluated by Applied Biosystems real-time PCR Instrument (ABI, Thermo Fisher, USA) in 10 μl reactions containing 2.5 μl SYBR Green PCR mix (TaKaRa, Japan) and 1 μl of each primer with 5 pmol/μl concentration. Specific human primers -including stemness related genes (OCT4, SOX2, NANOG, c-MYC and KLF4) and metastasis related genes (CDH1, CDH2, SNAIL1, TWIST1, TWIST2 and ZEB1) were used (Table S1) (See Supplementary Online Information at www.celljournal.org). PCR program was incubated at 95°C for 10 minutes, 40 cycles of denaturation at 95°C for 10 seconds, annealing at 60°C for 20 seconds and elongation at 72°C for 20 seconds. A final melting curve analysis from 65°C to 95°C was performed and the relative levels of expression were analyzed using 2^(-ΔΔCt) values. β-Actin was used as house-keeping gene.

**miRNA expression profiling**

miRNA expression levels were studied by performing SYBR Green qRT-PCR. In brief, 1 μg total RNA containing miRNAs was poly adenylated by poly (A) polymerase and reverse transcribed to cDNA using reverse transcriptase enzyme first strand cDNA synthesis reaction, provided from Parsgenome miR-Amp kit (Parsgenome, Iran).
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Iran) according to the manufacturer’s instructions. Each reaction was performed in a final volume of 10 μl, containing diluted cDNA and PCR master mix, and all reactions were run in triplicates. qRT-PCR reaction was performed using Applied Biosystems Real-Time PCR Instruments according to the manufacturer’s protocol. Expression levels of miRNA were normalized against internal controls U6, as a housekeeping control.

Statistical analysis

In vitro characterization of MCF-7 cell mammosphere and primary breast cancer tissue are presented as the mean ± SD of at least three different experiments. Two-tailed Student’s t test and analysis of variance (ANOVA) were performed to evaluate the difference between the mean values. The Spearman’s rank correlation test was used to evaluate miRNA and mRNA correlation. A two-tailed analysis with P<0.05 was considered statistically significant for all experiments.

For functional enrichment analysis, target genes of the selected miRNAs were submitted to Enrichr database. Subsequently, biological process, cellular component and molecular function were analyzed by Gene Ontology (GO) and pathways analysis was applied by KEGG 2017 (P<0.05).

Results

Computational analysis to identify common miRNA in stemness and EMT network

A total of 328 articles were yielded after the literature reviews, finally limited to 142 papers due to our exclusion criteria (mentioned in the method section). Full-text reviews were resulted in proposing 56 candidate miRNAs that have key role in BCSCs: 24 up-regulated and 32 down-regulated molecules. Among them, we chose miR-200c and miR-30c targeting at least three stemness and two EMT genes (Fig.1).

![Flowchart of studies represent systematic analysis to find miRNAs targeting both self-renewal and EMT pathways. EMT; Epithelial-mesenchymal transition, qRT-PCR; Quantitative real time polymerase chain reaction, GO; Gene ontology, KEGG; Kyoto Encyclopedia of genes and genomes-genomenet, GEO; Gene expression omnibus, and BSCs; Breast cancer stem cells.](image-url)
Mammospheres derived from MCF-7 as a model of breast cancer stem cells

MCF-7 cells were grown similar to adherent epithelial-like monolayer cells in culture (Fig.2A). Under serum-free and low-attachment conditions, MCF-7 cells grew into 3D non-linkage mammospheres within 24-48 hours, in comparison with their 2D adherent culture. Shape and appearance of spheres are solid and tightly packed in rounded margin, but we observed the mammospheres form looser and less rounded spheres over passages (Fig.2B-D). The secondary spheres were subsequently cultured up to three passages and MFE was calculated based on their size (> 60 μm, Fig.2E). The spheroid cells indicated about two folds increase (P<0.05) in MFE during three passages (Fig.2E). The results showed that MCF-7 cell-derived mammospheres had more colonogenic potential up to 5.5 fold. Indeed, the number as well as size of colonies was dominantly increased in mammospheres, in comparison with adherent cells (P<0.05, Fig.2F).

Mammospheres revealed increased ability of migration and invasion

The migratory capability of mammospheres was increased (about 1.7 fold). However, invasiveness potential of the isolated cells from mammospheres was significantly increased (about 3.35 fold), in comparison with adherent cells (Fig.2G, H).

Patients’ demography

Seven female breast cancer patients (mean age of 48 ± 8.04 years) were included in the study after signing written informed consent. The clinicopathological data of all patients has been shown in Table 1. All tumors were classified as invasive ductal carcinoma (IDC). Immunohistochemical study of ER and PR expressions were performed on surgical resection tissue samples of the tumors based on the standard methods. Three samples were positive for ER, PR and HER2 and four patients were undergoing neo-adjuvant therapy before sampling.

Expression of miR-200c-3p and miR-30c-5p in tumor/normal tissues and mammospheres/adherent cells

Findings showed that miR-200c was decreased in mammospheres, compared to parental MCF-7 cells (P=0.0025, Fig.3A, Right). Furthermore, this expression was down-regulated in breast cancers with metastatic conditions (patients I, II and V, Fig.3A, Left). In contrast, expression of miR-30c was overexpressed in mammospheres, compared to adherent cells (P=0.0011), and it was also up-regulated in three of patients with grade II/III who received neo-adjuvant therapy (Fig.3A).

Gene expression in tumor/normal tissues and mammospheres/adherent cells

In the next step, expression level of stemness related genes (OCT4, SOX2, KLF4, c-MYC and NANOG) and EMT transcription factors (CDH1, CDH2, SNAIL1, TWIST1, TWIST2 and ZEB1) were evaluated in all tumor samples and mammospheres. Interestingly, expression level of OCT4, SOX2 and c-MYC was significantly increased in mammospheres and three malignant breast tumors who were under neo-adjuvant therapy (patients I, II and V, Fig.3B, C). KLF4 was down-regulated in both tumor samples and mammospheres. Meanwhile, expression of NANOG was not changed in mammospheres, but it was down-regulated in tumors (Fig.3B). Among EMT-related genes, CDH2, SNAIL1, TWIST1/2 and ZEB1 were also overexpressed in mammospheres. However, tumors differentially expressed EMT related genes. Expression of CDH2, SNAIL1 and ZEB1 were up-regulated in three malignant breast tumors (patients I, II and V), but the others were down-regulated (Fig.3D, E). This demonstrates that signature of self-renewal related gene expressions and some EMT genes in malignant breast tumor of patients who underwent neo-adjuvant therapy are similar to that of mammospheres, as BCSC model.

| Patients | Age (Y) | Histological subtype | Ki-67 | Grade | ER status | PR status | HER2 status | Metastasis | Neo-adjuvant |
|----------|---------|----------------------|-------|-------|-----------|-----------|-------------|------------|-------------|
| Case I   | 56      | IDC                  | >10%  | III   | >80%      | >80%      | Negative    | Yes        | Yes         |
| Case II  | 51      | IDC                  | >30%  | III   | 60%       | 20%       | 30%         | Yes        | Yes         |
| Case III | 45      | IDC                  | NA    | I     | NA        | NA        | NA          | No         | Yes         |
| Case IV  | 50      | IDC                  | NA    | NA    | NA        | NA        | NA          | No         | No          |
| Case V   | 39      | IDC                  | >50%  | II    | >80%      | >80%      | >30%        | Yes        | Yes         |
| Case VI  | 58      | IDC                  | NA    | I     | NA        | NA        | NA          | No         | No          |
| Case VII | 37      | IDC                  | NA    | I     | NA        | NA        | NA          | No         | No          |

IDC; Invasive ductal carcinoma, ER; Estrogen receptor, PR; Progesterone receptor, and NA; Not available.

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Fig. 2: Colony and mammosphere formation abilities in MCF-7 and mammospheres. A. Parental cells cultured in 2D monolayer condition (magnification: ×4, scale bar: 100 μm). B-D. Serial mammospheres derived from the first generation of mammospheres (up to passage 3) showing progressive loss of cell cohesion and formed loose spheroid (magnification: ×20, scale bar: 100 μm). E. Mammosphere forming efficacy (MFE) was calculated from the first to third generation. Data are based on the mean percentages of the formed spheres quantity within a culture relative to the initial cell seeding number (mean ± SD, n=3). F. The percentage of colonies increased in cells derived from mammospheres in comparison to the adherent cells. (mean ± SD, n=3). G. Evaluation of migration and invasion abilities of the cells isolated from mammospheres and MCF-7 monolayer. Left panel shows crystal violet stained cells, passing through the matrigel coated filter insert (as invasive cells) or uncoated filter insert (as migratory cells), and H. Quantification of migratory and invasive cells in adherent vs. mammosphere cells. Mammospheres revealed higher migration and invasion rate than their parental cells. Bars indicated mean ± SD of three independent experiments. **; P<0.01 and ***; P<0.001.
**Fig. 3:** Expression level of miRNAs and genes. Expression levels of miR-200c and miR-30c as well as stemness and metastasis genes in human breast cancer versus normal breast (the first group), and mammospheres versus MCF-7 adherent cells (the second group) were determined by quantitative real time polymerase chain reaction (qRT-PCR). **A.** Expression of each miRNA was normalized to the levels of U6. Each cell line represents n≥3 and tumor represents n≥1. **B, C.** Scatter plot of stemness and metastasis gene expression levels in breast cancer and normal breast tissues (control). The line represents mean value, **D, and E.** Expression level of stemness and metastasis genes in mammospheres related to adherent cells (control), determined by qRT-PCR. β-Actin was used as the housekeeping gene. Statistically significant difference was determined by paired t test with GraphPad Prism 6 software. Bars indicated mean ± SEM.
Correlation of miR-200c and miR-30c with stemness and EMT genes, and with overall survival of breast invasive carcinoma

In next step, correlation of miR-200c and miR-30c with stemness and metastasis gene expressions were assessed in both mammospheres and tumor tissues. As shown in Table 2, miR-200c was negatively correlated to SOX2 and KLF4 stemness genes, as well as SNAIL and TWIST1 EMT genes. However, expression of miR-30c strongly displayed positive correlation with expression of most stemness related genes “OCT4, SOX2, KLF4 and NANOG” and all EMT related genes. Moreover, miR-200c had negative correlation with miR-30c (R=-0.8, P=0.04). All aforementioned data indicates discriminatory potential of miR-30c and miR-200c to target both EMT and self-renewal pathways in BCSCs and malignant breast tumors.

Target genes and pathways analyses for miR-200c-3p and miR-30c-5p

In order to recognize the potential miRNA efficacy for breast cancer tracing, we predicted target genes of miR-200c-3p and miR-30c-5p. They were listed to GO annotation dataset for analysis of molecular function, biological processes and cellular component by Enricher. The result was sorted based on p-value. The lowest P value is related to more specific term. GO analysis showed that targeted genes of the differentially expressed miRNAs were enriched in the molecular functions of E-box binding, DNA binding, N-box binding, estrogen response element binding, cadherin binding involved in cell-cell adhesion and miRNA binding (Fig.4A). The biological process of these genes included cell-cell adhesion, stem cell proliferation process, cell cycle, angiogenesis and EMT process (Fig.4B). In terms of cellular component, most of the genes belong to the nucleolus and cytoplasmic organelles (Fig.4C). Finally, KEGG pathway analysis also showed similar results, in terms of the number of genes involved in the adhesion junction, pathways in cancer, MAPK signaling pathway, Wnt signaling pathway, PI3K-AKT signaling pathway, HIF-1 signaling pathway, TGF-beta signaling pathway, as well as the signaling pathways regulating pluripotency of stem cells, P53 signaling pathway and cell cycle (Fig.4D). Additionally, using PROGmiR (14), we were able to create a significant diagnostic plot between the expression level of individual miR-200c and miR-30c, and overall survival rate of the patients. Actually, simultaneous deregulation of miR-200c and miR-30c could significantly reduce the survival rate of breast invasive carcinoma cells via up-regulation of OCT4, SOX2, c-MYC, SNAI1, ZEB1, CDH2 and down-regulation of CDH1 (P=0.02, Fig.4E).

### Table 2: Spearman’s rho for stemness and epithelial-mesenchymal transition (EMT) genes

| miR name | OCT4 | SOX2 | c-MYC | KLF4 | NANO | CDH1 | CDH2 | SNAIL1 | TWIST1 | TWIST2 | ZEB1 |
|----------|------|------|-------|------|------|------|------|--------|--------|--------|------|
| miR-200c | R=-0.4 | R=-0.94*** | R=-0.5 | R=-0.93*** | R=-0.4 | R=-0.9 | R=-0.4 | R=-0.6** | R=-0.8** | R=-0.4 | R=0.5 |
| P=0.4    | P<0.005 | P=0.08 | P=0.0006 | P=0.39 | P=0.01 | P=0.39 | P=0.01 | P=0.01 | P=0.32 | P=0.28 |
| miR-30c  | R=0.85*** | R=0.95*** | R=0.3  | R=0.71**  | R=0.9** | R=0.5 | R=0.74*** | R=0.87*** | R=0.92*** | R=0.82*** | R=0.83*** |
| P<0.000  | P=0.003 | P=0.29 | P=0.004 | P<0.001 | P=0.04 | P=0.002 | P=0.001 | P=2.9E-06 | P=0.0002 | P=0.0001 |

*; P<0.05, **; P<0.01, and ***; P<0.001.
Fig. 4: Gene ontology (GO), KEGG Pathway analysis using Enrichr and diagnostic plots creation with PROGmiR. **A.** Molecular function of stemness and epithelial-mesenchymal transition (EMT) regulated genes through the differentially expressed miRNAs, **B.** Biological process, **C.** Cellular component of these genes. Only the top ten enriched GO terms are represented, **D.** KEGG pathways with their P values. The most significant pathways bear the smallest P value listed from top to down, and **E.** Kaplan-Meier survival curve analysis was associated with overall survival in patients’ breast invasive carcinoma cells. The patients were stratified into high-risk and low-risk groups according to median of each miRNA.
Discussion

This study evaluates expression of miRNAs targeting both stemness and metastasis pathways in BCSCs. To provide BCSC model, we used MCF-7 derived mammospheres representing higher ability to sphere and colony formations, compared to their parental cells, as well as more invasion and migration capabilities. Our data are in agreement with the previous studies (15, 16) that observed MCF-7 derived mammospheres contain highest CSCs population (17, 18) with CD44+/CD24- phenotype (19, 20). Although, we did not evaluate tumorigenicity of the mammospheres in vivo, but adequate studies have determined tumorigenic ability of the cells originated from MCF-7-mammospheres in less than 1000 cell per injection (21).

To specify miRNAs involved in both stemness and metastasis regulation, systematic analysis was done using the important genes contributed to both pathways. The results implicated that has-miR-200c-3p and has-miR-30c-5p could potentially regulate these pathways. These miRNAs have been identified in different studies to control major transcription factors of EMT and induce metastasis (22, 23). It has also been reported that miR-200c controls BCSC functions (24, 25). Meanwhile, few studies implicated the role of miR-30c in BCSCs. Interestingly, the mammospheres of this study had similar expression pattern of miR-200c and miR-30c to three patients (I, II and V); these patients were at grade III/II and received neo-adjuvant therapy before sample collection. They showed significant down-regulation of miR-200c and up-regulation of miR-30c; however, miR-200c and miR-30c expression were both down-regulated in patients number six and four, among seven patients. Thus, by considering this similarity, we figured out the expression level of stemness related genes in mammospheres as well as all tissue samples. Impressively, expression of OCT4, SOX2 and c-MYC was up-regulated in mammospheres and the same three previous patients (I, II and IV). KLF4 expression, as another stemness related gene, was diminished in mammospheres and most of the tumor tissues, and NANOG was just significantly down-regulated in patient samples, but not in mammospheres. Moreover, transcription of miR-30c displayed positive correlation with OCT4, SOX2, KLF4 and NANOG expressions. In addition, miR-200c had negative correlation with expression of SOX2 and KLF4. Indeed, miR-200c significantly exhibited negative correlation with miR-30c. Similar to our data, miR-200c clusters (miR-200c-141, miR-200b-200a-429 and miR-183-96-182) have been reported to be down-regulated in isolated BCSCs from eleven human breast cancers tissues, normal mammary stem cells (26, 27) and carcinoma cells (28). Furthermore, lower expression of miR-200c in patients could be considered as a prognostic factor of breast cancer metastasis, since its down-regulation associates with poor survival rate. Up-regulation of this miRNA correlates with inhibition of cell proliferation and regulates cancer stem cell functions (29-31). In addition, miR-200c plays an important role in inhibiting proliferation of breast cancer cells by targeting the stemness related genes such as NANOG, SOX2 and KLF4 that are located in down-stream of miR-200c. It also inhibits tumor growth, differentiation and self-replication of CSCs by targeting TUBB3 and as a result it would be involved in restoring sensitivity to microtubule-targeting drugs (32).

In this study, miR-30c represents stronger correlation with most of the stemness related genes. This miRNA has previously been reported as a breast cancer prognostic biomarker (33) and its expression is various among different breast cancer subtypes. Higher miR-30c expression level was reported in luminal-A tumors and low miR-30c expression level was observed in basal-like tumors (34). In fact, few evidences are available representing effect of miR-30c in regulation of stemness, with mainly focus on EMT regulation. In one study, Yu et al. (35) showed down-regulation of miR-30 family, exclusively miR-30e, interferes with tumor beginning BCSCs (in mammospheres as well as primary BCSCs acquired from breast cancer patients) through up-regulation of ubiquitin-conjugating enzyme 9 (Ubc9) and integrin β3 (ITGB3). This up-regulation results in reduced self-renewal and anti-apoptotic features of BCSCs. Overexpression of miR-30a considerably decreased the sphere creation capability of MCF-7 cells, while derepression of miR-30a intensely enhanced the number of mammospheres in the human breast cancer cell line, MCF-7 (36).

Consistent with other studies, the present study demonstrated correlation of miR-200c and miR-30c with expression of important EMT transcription factors (SNAIL, TWIST and ZEB1), in tumors and mammospheres. miR-200c not only is a malignancy biomarker, but also promote metastasis in poor metastatic cells in vivo, presence of which in serum of metastatic breast cancer patients can be indicated for brain metastases (37). miR-200c maintains cells in an epithelial state condition, via the regulation of mesenchymal genes such as CDH2, SNAI1, SNAI2, TWIST1, TWIST2 and ZEB1 (27). miR-30c contributes to miRNA-cytoskeleton regulation network and its target genes (i.e. VIM, TWF1, and IL-11) represent invasion, EMT and chemo-resistance molecular mechanisms (35). In treatment of breast cancer cells, miR-200c was also reported to induce apoptosis (38) and sensitize the cells to chemotherapy, radiotherapy and trastuzumab using therapy. Down-regulation of this molecule is known as marker for drug resistance in female genital tumors, such as ovarian, cervical and breast cancers (39).

We further employed bioinformatics tools to find out the target genes and pathways of miR-200c and miR-30c coordinating stemness and metastasis. Pathway analysis indicated that these genes considerably associate with "adherens junction pathway", "pathways involved in cancer", "MAPK signaling pathway", "Wnt signaling pathway", "PI3K-Akt signaling pathway", "regulating pluripotency of stem cells", "P53 signaling pathway", "TGFβ signaling pathway" and "HIF-1 signaling".
All of these pathways have been reported to be related to several cellular activities including proliferation, migration, invasion, cell cycle, regulation of ER signaling in cancer and CSCs. Adherens junction pathways are also of the major mechanisms presented in stem cells, where raising documents have illustrated that remodeling of the cytoskeletal proteins could characterize stem cell destiny (40). To the best of our knowledge, this is the first experiment comparing mammmospheres as BSCSs model with signature pattern of metastatic patients (pre or post neo-adjuvant therapy). Because of similarity of miR-200c and miR-30c expression levels in mammmospheres and some of our patients (I, II and V), we suggest that combination of these miRNAs might predict outcome of adjuvant therapy or metastasis in patients. Down-regulation of miR-200c and up-regulation of miR-30c suggest that metastatic breast tumors and mammmospheres are similar and they contribute to communal molecular mechanisms regulating stem cell functions such as self-renewal, proliferation, EMT and resistance to drug.

Conclusion

The present study demonstrates that down-regulation of miR-200c and up-regulation of miR-30c promote BSCS features toward malignant breast tumors, leading to their resistance to neo-adjuvant therapy. These findings suggest a signature to predict metastasis post chemotherapy in breast cancer patients. However, further experiments are required in this regard.

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Authors’ Contributions

M.E.; Participated in study design, interpretation of data, drafting and statistical analysis. E.A.; Contributed extensively in interpretation of the data, drafting and statistical analysis and the conclusion. M.R.; Contributed to all experimental work, data collection and evaluation and statistical analysis. A.Sh.-Z.; Conducted statistical and bioinformatics analysis. N.Z.; Participated in study design, drafting and approved the final version of this manuscript for submission. L.G.; Performed sample collection and prepared breast cancer tumor for molecular analysis to this component of the manuscript. All authors read and approved the final manuscript.

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