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## Data Article

# Dataset for the proteomic inventory and quantitative analysis of the breast cancer hypoxic secretome associated with osteotropism



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## ABSTRACT

The cancer secretome includes all of the macromolecules secreted by cells into their microenvironment. Cancer cell secretomes are significantly different to that of normal cells reflecting the changes that normal cells have undergone during their transition to malignancy. More importantly, cancer secretomes are known to be active mediators of both local and distant host cells and play an important role in the progression and dissemination of cancer. Here we have quantitatively profiled both the composition of breast cancer secretomes associated with osteotropism, and their modulation under normoxic and hypoxic conditions. We detect and quantify 162 secretome proteins across all conditions which show differential hypoxic induction and association with osteotropism. Mass Spectrometry proteomics data have been deposited to the ProteomeXchange Consortium with the dataset identifier PXD000397 and the complete proteomic, bioinformatic and biological analyses are reported in Cox et al. (2015) [1].

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## Specifications table

Subject area	Biology
More specific sub-ject area	Breast Cancer, Bone Metastasis, Secretome
Type of data	Mass Spectrometry RAW files
How data was acquired	LC-MS/MS on a Q-Exactive Mass Spectrometer (Thermo Fisher Scientific)
Data format	.RAW files
Experimental factors	None applied
Experimental features	The project profiled the expression patterns in hypoxia induced secretomes between MDA-MB-231 parental and MDA-MB-231 Bone Tropic (BT) breast cancer cell lines which have been previously generated by Massague and colleagues (Kang et al. Cancer Cell 2003).
Data source location	Copenhagen, Denmark
Data accessibility	The Mass Spectrometry proteomics data for the article “The hypoxic cancer secretome induces pre-metastatic bone lesions through lysyl oxidase” doi:10.1038/nature14492 [1] have been deposited to the ProteomeXchange Consortium ( <a href="http://proteomecentral.proteomexchange.org">http://proteomecentral.proteomexchange.org</a> ) via the PRIDE partner repository [2] with the dataset identifier PXD000397

## Value of the data

- This data set will be of value to the scientific community wanting to determine which proteins are secreted from breast cancer cells that metastasize to bone, and those that are regulated by hypoxia.
- The dataset includes quantitative global proteome (Label-free and SILAC) analysis of hypoxic induced secretomes of parental and bone tropic human breast cancer cell lines.
- Differential analysis of secretome changes associated with bone tropism in breast cancer.
- Response of the cancer secretome to hypoxic induction.

## 1. Data, experimental design, materials and methods

### 1.1. Cell lines

The MDA-MB-231 Bone Tropic (BT) 1833 subclone cell line was obtained from J. Massagué at the Memorial Sloan-Kettering Cancer Center. The MDA-MB-231 parental cell line was obtained from the American Type Culture Collection (ATCC) (distributed by LGC Standards). All cell lines were routinely tested for mycoplasma and tested negative for murine pathogens by IMPACT I testing (IDEXX Laboratories).

### 1.2. Collection of hypoxia induced conditioned medium (CM)

For label-free Mass Spectrometry analysis, the MDA-MB-231 parental (wt) cell line and the MDA-MB-231 Bone Tropic (BT) (1833 sub clone derived in Kang et al. [3]) (a kind gift from J. Massagué) were routinely cultured in high glucose Dulbecco's Modified Essential Media (DMEM)+ GlutaMAX (Gibco 31966-021) with 100 U/ml penicillin and 100 µg/ml streptomycin (Gibco 15140-122), plus 10% dialysed Fetal Bovine Serum (FBS) (Gibco 26400). For conditioned media (CM) collection, cells were

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